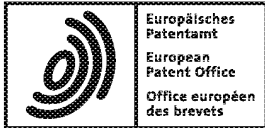


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(54) **Ligands for modulating the expression of exogenous genes via an ecdysone receptor complex**

Liganden zur Beeinflussung der Expression exogener Gene mittels eines Ecdysone-Rezeptor-Komplexes

Ligands qui module l'expression des genes exogenes à travers du complexe de recepteur d'ecdysone

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Description

[0001] This invention relates to the use of non-steroidal ligands for inducing or suppressing the expression of an exogenous gene in animal and plant cells.

[0002] In the field of genetic engineering, precise temporal control of gene expression, that is, the ability to activate or suppress a gene, is a valuable tool in studying, manipulating, and controlling development and other physiological processes (see, for example, Evans and No, PCT International Publication No. WO97/038117 and references cited therein). In mammalian systems, applications include inducible gene targeting, overexpression of toxic and teratogenic genes, anti-sense RNA expression, and gene therapy. In plant systems, applications include the control of plant traits, male or female fertility; overexpression of plant protective agents; and production of or modification of desired plant products including both native and non-native materials. For both animals and plants, inducibility can be valuable for foreign protein production, for example, therapeutic proteins, industrial enzymes, polymers, and the like.

[0003] It is important that the agent used to control gene expression, which is often referred to as a "gene switch", be one which is normally absent from the organism in which the gene to be controlled resides. This is to avoid unexpected expression or suppression of the gene. For example, an inducible tetracycline regulated system has been devised and utilized in transgenic mice whereby gene activity is induced in the absence of the antibiotic and suppressed in its presence. Unfortunately, in this case, the pharmacokinetics of tetracycline may interfere with its use as an efficient "on-off" gene switch.

[0004] International Patent Publication No. WO 96/037609 describes an insect steroid receptor isolated from *Heliothis virescens* ("HEcR") which is capable of acting as a gene switch responsive to both steroid (e.g., 20-hydroxyecdysone and Muristerone A) and certain non-steroidal inducers. Non-steroidal inducers have a distinct advantage over steroids, in this and many other systems which are responsive to both steroids and non-steroid inducers, for a number of reasons including, for example: lower manufacturing cost, metabolic stability, absence from insects, plants, or mammals, and environmental acceptability. The PCT application describes the utility of two dibenzoylhydrazines, 1,2-dibenzoyl-1-*tert*-butyl-hydrazine and tebufenozide (N-(4-ethylbenzoyl)-N'-(3,3-dimethylbenzoyl)-N'-*tert*-butyl-hydrazine) as gene switches for the HEcR system and suggests that other dibenzoylhydrazines, such as those disclosed in U.S. Patent No 5,117,057 may also function as gene switches in the system. While this may be true, activity of these dibenzoylhydrazines is uncertain. Specifically, 5,117,057 shows a very broad class of dibenzoylhydrazines, many of which appear to be ineffective as gene switches. WO 96/037609 indicates that when 20 such dibenzoylhydrazines were tested, only 7 showed any activity.

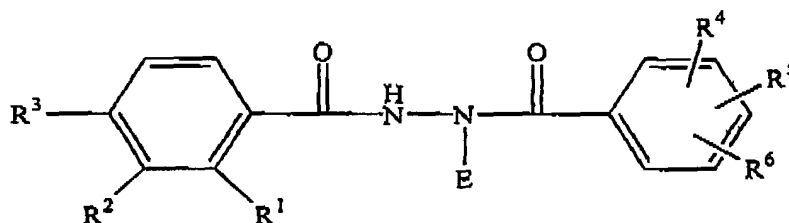
[0005] International Patent Publication No. WO 96/027673 discloses the use of tebufenozide as a chemical ligand for the ecdysone receptor from *Drosophila melanogaster*. This receptor is used to control gene expression in transgenic plants resulting in the control of various traits of agronomic importance.

[0006] Unfortunately, even though the ligands described in the above-identified references show reporter gene induction activity in isolated cells, no consideration was made for their use in whole organisms such as intact plants, and animals.

[0007] Therefore, there remains a continuing need to develop non-steroidal ligands with increased or consistent activity compared to known ligands and which demonstrate that activity in intact plants and animals. We have discovered a limited group of dibenzoylhydrazine derivatives which not only show reporter gene induction activity in isolated cells but also have advantages over the known diacylhydrazines, 1,2-dibenzoyl-1-*tert*-butyl-hydrazine and tebufenozide, when used in intact plants and animals, due to their improved transport and distribution properties, metabolic stability, residual activity, affinity for the receptor, and lack of adverse effects.

[0008] In one aspect of the present invention there is provided an *in vitro* method to modulate exogenous gene expression comprising contacting an ecdysone receptor complex comprising:

- a) a DNA binding domain;
- b) a ligand binding domain;
- c) a transactivation domain; and
- d) a ligand of the formula:



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt;

provided that:

a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl ;

c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is:

i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or

ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, CN, OMe, or OEt;

d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;

then R¹ is Me or Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SOMe; then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

with a DNA construct comprising:

a) the exogenous gene; and

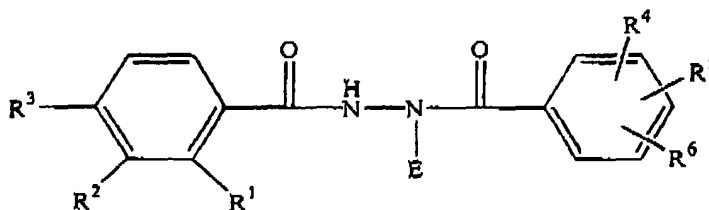
b) a response element;

wherein:

a) the exogenous gene is under the control of the response element; and

b) binding of the DNA, binding domain to the response element in the presence of the ligand results in activation or suppression of the gene,

[0009] In another aspect, the present invention provides a method to modulate the expression of one or more exogenous genes in a plant, comprising administering to the plant an effective amount of a ligand of the formula:



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt;

provided that:

a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl ;

c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is:

i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or

ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, CN, OMe, or OEt;

d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SOMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

wherein the cells of the plant contain:

a) an ecdysone receptor complex comprising:

1) a DNA binding domain;

2) a binding domain for the ligand; and

3) a transactivation domain; and

b) a DNA construct comprising:

1) the exogenous gene; and

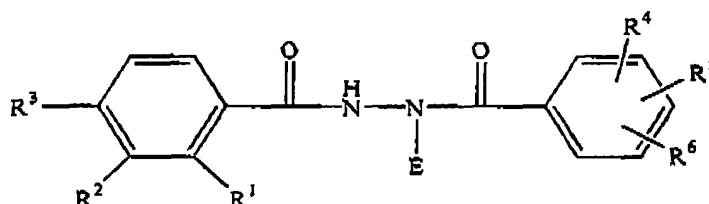
2) a response element; and

wherein:

- a) the exogenous gene is under the control of the response element; and
- b) binding of the DNA binding domain to the response element in the presence of the ligand results in activation or suppression of the gene.

[0010] There is provided in a further aspect of the present invention a method for producing a polypeptide comprising the steps of:

- a) selecting a cell which is insensitive to exposure to a ligand of the formula:



wherein;

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt;

provided that:

- a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

- b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl ;

- c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is:

- i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or
- ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, OMe, or OEt;

- d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

- e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

- f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

- g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

- h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

- b) introducing into the cell:

1) a DNA construct comprising:

- a) an exogenous gene encoding the polypeptide; and
- b) a response element;

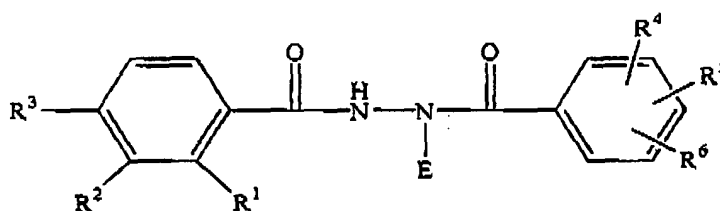
wherein the gene is under the control of the response element; and

2) an ecdysone receptor complex comprising:

- a) a DNA binding domain;
- b) a binding domain for the ligand; and
- c) a transactivation domain; and

c) exposing the cell to the ligand of step(a).

[0011] In another aspect of the present invention there is provided a method for regulating endogenous or heterologous gene expression in a transgenic plant comprising contacting a ligand of the formula:



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt;

provided that:

a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl;

c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is:

i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or

ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, CN, OMe, or OEt;

d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl

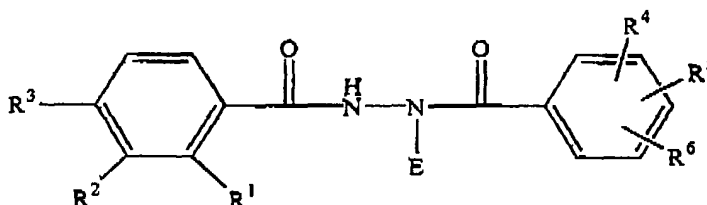
ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SOMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; with an ecdysone receptor complex within the cells of the plant, wherein the cells further contain a DNA binding sequence for the ecdysone receptor complex when in combination with the ligand and wherein formation of an ecdysone receptor complex-ligand-DNA binding sequence complex induces expression of the gene.

[0012] In a further aspect the present invention provides the use of a ligand of the formula (I):



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOME, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt;

provided that:

a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl;

c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is;

i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or

ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, CN, OMe, or OEt;

d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SOMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

for modulating *in vitro* the expression of one or more exogenous genes in an animal cell, wherein the cell contains:

a) an ecdysone receptor complex comprising:

- 1) a DNA binding domain;
- 2) a binding domain for the ligand; and
- 3) a transactivation domain; and

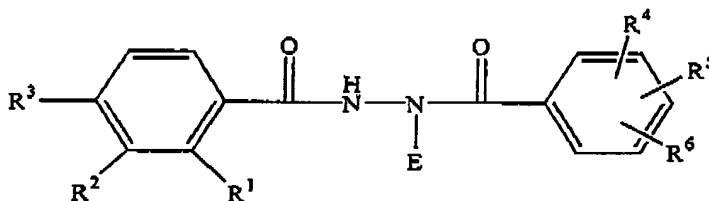
b) a DNA construct comprising:

- 1) the exogenous gene; and
- 2) a response element; and

wherein:

- a) the exogenous gene is under the control of the response element; and
- b) binding of the DNA binding domain to the response element in the presence of the ligand results in activation or suppression of the gene.

[0013] In another aspect of the present invention, there is provided the use of a ligand of the formula (I):



wherein:

E is a $(\text{C}_4\text{-C}_6)$ alkyl containing a tertiary carbon;

R^1 is H, Me, Et, i-Pr, F, or OMe;

R^2 is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe, or joined with R^3 and the phenyl carbons to which R^2 and R^3 are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R^3 is H, Et, or joined with R^2 and the phenyl carbons to which R^2 and R^3 are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R^4 , R^5 , and R^6 are independently H, Me, Et, F, Cl, CH_2OH , CN, OMe, or OEt;

provided that:

a) when R^1 is Me and R^2 is OMe;

then R^3 is H; and the combination R^4 , R^5 , and R^6 is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R^1 is Me and R^2 is OEt;

then R^3 is H and the combination R^4 , R^5 , and R^6 is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl ;

c) when R^1 is Et and R^2 is OMe or OEt;

then R^3 is H and the combination R^4 , R^5 , and R^6 is:

- i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or
- ii) R^6 is H, R^4 is Me, and R^5 is Et, F, Cl, CH_2OH , CN, OMe, or OEt;

d) when R^1 is i-Pr;

then R^2 is OMe, or OEt; R^3 is H; and the combination R^4 , R^5 , and R^6 is 3,5-di-Me;

e) when R^3 is Et;

then R^2 is H, R^1 is F, and the combination R^4 , R^5 , and R^6 is 3,5-di-Me;

- f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;
 then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;
 g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl
 ring;
 then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and
 h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SOMe;
 then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

in the preparation of a pharmaceutical for modulating the expression of one or more exogenous genes in an animal,
 wherein cells of the animal contain:

a) an ecdysone receptor complex comprising:

- 1) a DNA binding domain;
- 2) a binding domain for the ligand; and
- 3) a transactivation domain; and

b) a DNA construct comprising:

- 1) the exogenous gene; and
- 2) a response element; and

wherein:

- a) the exogenous gene is under the control of the response element; and
- b) binding of the DNA binding domain to the response element in the presence of the ligand results in activation or suppression of the gene.

[0014] There is described herein an improvement in a method to modulate exogenous gene expression comprising
 contacting an ecdysone receptor complex comprising:

- a) a DNA binding domain;
- b) a ligand binding domain;
- c) a transactivation domain; and
- d) a ligand;

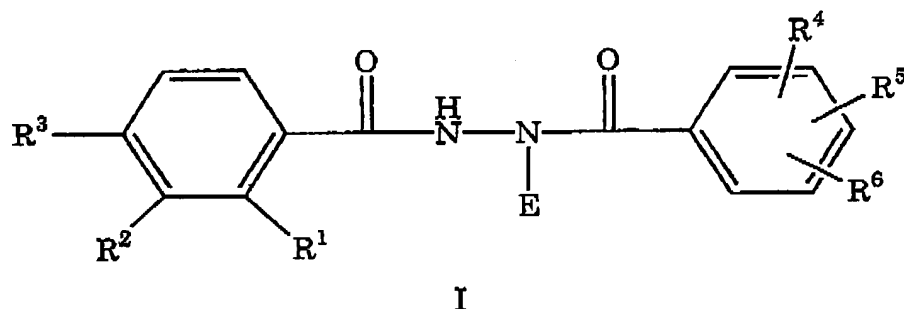
with a DNA construct comprising:

- a) the exogenous gene; and
- b) a response element;

wherein:

- a) the exogenous gene is under the control of the response element; and
- b) binding of the DNA binding domain to the response element in the presence of the ligand results in activation or suppression of the gene; the improvement comprising:

selecting the ligand from a compound of formula I:



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon or a cyano(C₃-C₆)alkyl containing a tertiary carbon;
 R¹ is H, Me, Et, i-Pr, F, formyl, CF₃, CHF₂, CHCl₂, CH₂F, CH₂Cl, CH₂OH, CH₂OMe, CH₂CN, CN, C≡CH, 1-propynyl, 2-propynyl, vinyl, OH, OMe, OEt, cyclopropyl, CF₂CF₃, CH=CHCN, allyl, azido, SCN, or SCHF₂;
 R² is H, Me, Et, n-Pr, i-Pr, formyl, CF₃, CHF₂, CHCl₂, CH₂F, CH₂Cl, CH₂OH, CH₂OMe, CH₂CN, CN, C≡CH, 1-propynyl, 2-propynyl, vinyl, Ac, F, Cl, OH, OMe, OEt, O-n-Pr, OAc, NMe₂, NEt₂, SMe, SEt, SOCF₃, OCF₂CF₂H, COEt, cyclopropyl, CF₂CF₃, CH=CHCN, allyl, azido, OCF₃, OCHF₂, O-i-Pr, SCN, SCHF₂, SMe, NH-CN, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;
 R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;
 R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, Br, formyl, CF₃, CHF₂, CHCl₂, CH₂F, CH₂Cl, CH₂OH, CN, C≡CH, 1-propynyl, 2-propynyl, vinyl, OMe, OEt, SMe, or SEt;

provided that:

- a) when R¹ is Me and R² is OMe;
 then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;
- b) when R¹ is Me and R² is OEt;
 then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl ;
- c) when R¹ is Et and R² is OMe or OEt;
 then R³ is H and the combination R⁴, R⁵, and R⁶ is:
 - i) 3, 5-di-OMe-4-Me, 3, 5-di-Cl, 3, 5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Br-5-Me, 2-Cl, 2-Br, or 3-Me; or
 - ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, Br, formyl, CF₃, CHF₂, CHCl₂, CH₂F, CH₂Cl, CH₂OH, CN, C≡CH, 1-propynyl, 2-propynyl, vinyl, OMe, OEt, SMe, or SEt;
- d) when R¹ is i-Pr;
 then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;
- e) when R³ is Et;
 then R² is H, R¹ is F or Cl, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;
- f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;
 then R¹ is Me or Et and the combination R⁴, R⁵, and R⁶ is 3, 5-di-Me;
- g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;
 then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;
- h) when R¹ is formyl, CF₃, CHF₂, CHCl₂, CH₂F, CH₂Cl, CH₂OH, CH₂OMe, CH₂CN, CN, C≡CH, 1-propynyl, 2-propynyl, vinyl, OH, cyclopropyl, CF₂CF₃, CH=CHCN, allyl, azido, SCN, or SCHF₂;
 then R² is OMe or OEt, R³ is H, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and
- i) when R² is Me, Et, n-Pr, i-Pr, formyl, CF₃, CHF₂, CHCl₂, CH₂F, CH₂Cl, CH₂OH, CH₂OMe, CH₂CN, CN, C≡CH,

1-propynyl, 2-propynyl, vinyl, Ac, F, Cl, OH, O-n-Pr, OAc, NMe₂, NEt₂, SMe, SEt, SOCF₃, OCF₂CF₂H, COEt, cyclopropyl, CF₂CF₃, CH=CHCN, allyl, azido, OCF₃, OCHF₂, O-i-Pr, SCN, SCHF₂, SMe, or NH-CN; then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me.

[0015] There is mentioned herein a method to modulate exogenous gene expression comprising contacting an ecdysone receptor complex comprising:

- a) a DNA binding domain;
- b) a ligand binding domain;
- c) a transactivation domain; and
- d) a ligand consisting of a compound of formula I:

with a DNA construct comprising:

- a) the exogenous gene; and
- b) a response element;

wherein:

- a) the exogenous gene is under the control of the response element; and
- b) binding of the DNA binding domain to the response element in the presence of the ligand results in activation or suppression of the gene.

[0016] In order to achieve an optimum balance between a) ligand binding and the resulting gene switch activity, and b) transport, systemicity, toxicity, and metabolic stability in intact plants and animals, the position and size of the compound of formula I substituent groups are important. The optimum balance of properties appears to occur when E is t-butyl, R¹ is ethyl, R² is ethoxy, R³ is hydrogen, and the combination R⁴, R⁵, and R⁶ is 3,5-dimethyl. The composition of each of these "R" groups may be varied considerably. However, variations which lead to significant changes in the size, shape, and overall polarity of the compound of formula I will tend to reduce the optimum balance and, consequently, the improved properties of the compound. For this reason, when the composition of any one particular R group is changed from the optimum, variations in the composition of the remaining R groups should be limited.

[0017] Preferably, E is (C₄-C₅)alkyl. More preferably, E is t-butyl.

[0018] Preferably, R¹ is Me, Et, i-Pr, F, CF₃, CHF₂, CHCl₂, CH₂F, CH₂Cl, CH₂OH, CH₂OMe, CH₂CN, CN, C≡CH, or CF₂CF₃. More preferably, R¹ is Me, Et, i-Pr, F, CF₃, CHF₂, CH₂F, CH₂OMe, CH₂CN, C≡CH, or CF₂CF₃. Even more preferably, R¹ is Me, Et, i-Pr, or F. Most preferably, R¹ is Me or Et.

[0019] Preferably, R² is Et, CF₃, CHF₂, CHCl₂, CH₂F, CH₂Cl, CH₂OH, CH₂OMe, CH₂CN, CN, OH, OMe, OEt, O-n-Pr, CF₂CF₃, azido, OCF₃, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon. More preferably, R² is Et, CF₃, CHF₂, CHCl₂, CH₂F, CH₂Cl, CH₂OH, CH₂OMe, CH₂CN, CN, OH, OMe, OEt, CF₂CF₃, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy or dihydrofuryl ring with the oxygen adjacent to a phenyl carbon. Even more preferably, R² is OH, OMe, OEt, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy or dihydrofuryl ring with the oxygen adjacent to a phenyl carbon. Most preferably, R² is OMe or OEt.

[0020] Preferably, R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon. More preferably, R³ is H, Et or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy or dihydrofuryl ring with the oxygen adjacent to a phenyl carbon. Even more preferably, R³ is joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy or dihydrofuryl ring with the oxygen adjacent to a phenyl carbon.

[0021] Preferably, R⁴, R⁵, and R⁶ are independently Me, F, Cl, CH₂OH, or OMe. More preferably, R⁴, R⁵, and R⁶ are independently Me, F, Cl, CH₂OH, or OMe. Even more preferably, R⁴, R⁵, and R⁶ are independently Me, F, or Cl. Still more preferably, the combination of R⁴, R⁵, and R⁶ is 3,5-di-Me, 3, 5-di-Cl, or 3, 5-di-F. Most preferably, the combination of R⁴, R⁵, and R⁶ is 3,5-di-Me.

[0022] The terms "Me", "Et", "n-Pr", "i-Pr", and "Ac" mean methyl, ethyl, normal propyl, isopropyl, and acetyl, respectively. When referring to R⁴, R⁵, and R⁶, the term "2,4", "2,5", "3,5", and the like refer to the relative positions on the phenyl ring to which the groups are attached.

[0023] The term "halo" means fluoro, chloro, bromo, or iodo.

[0024] The term "modulate" means the ability of a given ligand/receptor complex to induce or suppress the transac-

tivation of an exogenous gene.

[0025] The term "exogenous gene" means a gene foreign to the subject, that is, a gene which is introduced into the subject through a transformation process or an unmutated version of an endogenous mutated gene. The method of transformation is not critical and may be any method suitable for the subject known to those in the art. For example, transgenic plants are obtained by regeneration from the transformed cells. Numerous transformation procedures are known from the literature such as agroinfection using *Agrobacterium tumefaciens* or its T₁ plasmid, electroporation, microinjection of plant cells and protoplasts, and microprojectile transformation. Complementary techniques are known for transformation of animal cells and regeneration of such transformed cells in transgenic animals. Exogenous genes can be either natural or synthetic genes and therapeutic genes which are introduced into the subject in the form of DNA or RNA which may function through a DNA intermediate such as by reverse transcriptase. Such genes can be introduced into target cells, directly introduced into the subject, or indirectly introduced by the transfer of transformed cells into the subject. The term "therapeutic gene" means a gene which imparts a beneficial function to the host cell in which such gene is expressed. Therapeutic genes are not naturally found in host cells.

[0026] The term "ecdysone receptor complex" generally refers to a heterodimeric protein complex consisting of two members of the steroid receptor family, ecdysone receptor ("EcR") and ultraspiracle ("USP") proteins (see Yao, T.P., et. al. (1993) Nature 366, 476-479; Yao, T.-P., et. al., (1992) Cell 71, 63-72). The functional ecdysteroid receptor complex may also include additional protein(s) such as immunophilins. Additional members of the steroid receptor family of proteins, known as transcriptional factors (such as DHR38, *betaFTZ-1* or other insect homologs), may also be ligand dependent or independent partners for EcR and/or USP. The ecdysone receptor complex can also be a heterodimer of ecdysone receptor protein and the vertebrate homolog of ultraspiracle protein, retinoic acid-X-receptor ("RXR") protein. Homodimer complexes of the ecdysone receptor protein or USP may also be functional under some circumstances.

[0027] An ecdysteroid receptor complex can be activated by an active ecdysteroid or non-steroidal ligand bound to one of the proteins of the complex, inclusive of EcR, but not excluding other proteins of the complex.

[0028] The ecdysone receptor complex includes proteins which are members of the steroid receptor superfamily wherein all members are characterized by the presence of an amino-terminal transactivation domain, a DNA binding domain ("DBD"), and a ligand binding domain ("LBD") separated by a hinge region. Some members of the family may also have another transactivation domain on the carboxy-terminal side of the LBD. The DBD is characterized by the presence of two cysteine zinc fingers between which are two amino acid motifs, the P-box and the D-box, which confer specificity for ecdysone response elements. These domains may be either native, modified, or chimeras of different domains of heterologous receptor proteins.

[0029] The term "response element" ("RE") means one or more cis-acting DNA elements which confer responsiveness on a promoter mediated through interaction with the DNA-binding domains of the ecdysone receptor complex. This DNA element may be either palindromic (perfect or imperfect) in its sequence or composed of sequence motifs or half sites separated by a variable number of nucleotides. The half sites can be similar or identical and arranged as either direct or inverted repeats. The ecdysone receptor complex binds, in the presence or absence of a ligand, to the DNA sequence of an RE to initiate or suppress transcription of downstream gene(s) under the regulation of this response element. Examples of DNA sequences for RE of the natural ecdysone receptor include: RRGG/TTCANTGAC/ACYY (see Cherbas L., et. al., (1991), Genes Dev. 5, 120-131); AGGTCAN_(n)AGGTCA, where N_(n) can be one or more spacer nucleotides (see D'Avino PP., et. al., (1995), Mol. Cell. Endocrinol. 113, 1-9); and GGGTTGAATGAATTT (see Antoniewski C., et. al., (1994), Mol. Cell Biol. 14, 4465-4474).

[0030] The DNA sequences making up the exogenous gene, the response element, and the ecdysone receptor complex may be incorporated into procaryotic cells such as *Escherichia coli*, *Bacillus subtilis*, or other enterobacteria. However, because many of the proteins expressed by the gene are processed incorrectly in bacteria, eucaryotic cells are preferred. The nucleotide sequences for the exogenous gene, the response element, and the receptor complex can also be incorporated as RNA molecules, preferably in the form of functional viral RNAs such as tobacco mosaic virus. Of the eucaryotic cells, vertebrate cells are preferred because they naturally lack the molecules which confer responses to the ligands for the ecdysone receptor. As a result, they are insensitive to the ligands described herein. Thus, the ligands described herein will have negligible physiological or other effects on transformed cells, or the whole organism. Therefore, cells can grow and express the desired product, substantially unaffected by the presence of the ligand itself.

[0031] The term "subject" means an intact plant or animal or a cell from a plant or animal. It is also anticipated that the ligands will work equally well when the subject is a fungus or yeast. When the subject is an intact animal, preferably the animal is a vertebrate, most preferably a mammal.

[0032] The ligands described herein, when used with the ecdysone receptor complex which in turn is bound to the response element linked to an exogenous gene, provide the means for external temporal regulation of expression of the exogenous gene. The order in which the various components bind to each other, that is, ligand to receptor complex and receptor complex to response element, is not critical. Typically, modulation of expression of the exogenous gene is in response to the binding of the ecdysone receptor complex to a specific control, or regulatory, DNA element. The ecdysone receptor protein, like other members of the steroid receptor family, possesses at least three domains, a

transactivation domain, a DNA binding domain, and a ligand binding domain. This receptor, like a subset of the steroid receptor family, also possesses less well defined regions responsible for heterodimerization properties. Binding of the ligand to the ligand binding domain of ecdysone receptor protein, after heterodimerization with USP or RXR protein, enables the DNA binding domains of the heterodimeric proteins to bind to the response element in an activated form, thus resulting in expression or suppression of the exogenous gene. This mechanism does not exclude the potential for ligand binding to either EcR or USP, and the resulting formation of active homodimer complexes (e.g. EcR+EcR or USP+USP). Preferably, one or more of the receptor domains can be varied producing a chimeric gene switch. Typically, one or more of the three domains may be chosen from a source different than the source of the other domains so that the chimeric receptor is optimized in the chosen host cell or organism for transactivating activity, complementary binding of the ligand, and recognition of a specific response element. In addition, the response element itself can be modified or substituted with response elements for other DNA binding protein domains such as the GAL-4 protein from yeast (see Sadowski, et al. (1988) Nature, 335, 563-564) or LexA protein from *E. coli* (see Brent and Ptashne (1985), Cell, 43, 729-736) to accommodate chimeric ecdysone receptor complexes. Another advantage of chimeric systems is that they allow choice of a promoter used to drive the exogenous gene according to a desired end result. Such double control can be particularly important in areas of gene therapy, especially when cytotoxic proteins are produced, because both the timing of expression as well as the cells wherein expression occurs can be controlled. The term "promoter" means a specific nucleotide sequence recognized by RNA polymerase. The sequence is the site at which transcription can be specifically initiated under proper conditions. When exogenous genes, operatively linked to a suitable promoter, are introduced into the cells of the subject, expression of the exogenous genes is controlled by the presence of the ligand described herein. Promoters may be constitutively or inducibly regulated or may be tissue-specific (that is, expressed only in a particular type of cells) or specific to certain developmental stages of the organism.

[0033] Another aspect described herein is a method to modulate the expression of one or more exogenous genes in an organism, comprising administering to the organism an effective amount, that is, the amount required to elicit the desired gene expression or suppression, of a ligand comprising a compound of formula I and wherein the cells of the organism contain:

a) an ecdysone receptor complex comprising:

- 1) a DNA, binding domain;
- 2) a binding domain for the ligand; and
- 3) a transactivation domain; and

b) a DNA construct comprising:

- 1) the exogenous gene; and
- 2) a response element; and

wherein:

- a) the exogenous gene is under the control of the response element; and
- b) binding of the DNA binding domain to the response element in the presence of the ligand results in activation or suppression of the gene.

[0034] A related aspect described herein is a method for regulating endogenous or heterologous gene expression in a transgenic organism comprising contacting a ligand comprising a compound of formula I with an ecdysone receptor within the cells of the organism wherein the cells contain a DNA binding sequence for the ecdysone receptor and wherein formation of an ecdysone receptor-ligand-DNA binding sequence complex induces expression of the gene.

[0035] A fourth aspect described herein is a method for producing a polypeptide comprising the steps of:

- a) selecting a cell which is substantially insensitive to exposure to a ligand comprising a compound of formula I;
- b) introducing into the cell:

1) a DNA construct comprising:

- a) an exogenous gene encoding the polypeptide; and
- b) a response element;

wherein the gene is under the control of the response element; and

2) an ecdysone receptor complex comprising:

- a) a DNA binding domain;
- b) a binding domain for the ligand; and
- c) a transactivation domain; and

c) exposing the cell to the ligand.

[0036] As well as the advantage of temporally controlling polypeptide production by the cell, this aspect provides a further advantage, in those cases when accumulation of such a polypeptide can damage the cell, in that expression of the polypeptide may be limited to short periods. Such control is particularly important when the exogenous gene is a therapeutic gene. Therapeutic genes may be called upon to produce polypeptides which control needed functions, such as the production of insulin in diabetic patients. They may also be used to produce damaging or even lethal proteins, such as those lethal to cancer cells. Such control may also be important when the protein levels produced may constitute a metabolic drain on growth or reproduction, such as in transgenic plants.

[0037] Numerous genomic and cDNA nucleic acid sequences coding for a variety of polypeptides are well known in the art. Exogenous genetic material useful with the ligands described herein include genes that encode biologically active proteins of interest, such as, for example, secretory proteins that can be released from a cell; enzymes that can metabolize a substrate from a toxic substance to a non-toxic substance, or from an inactive substance to an active substance; regulatory proteins; cell surface receptors; and the like. Useful genes also include genes that encode blood clotting factors, hormones such as insulin, parathyroid hormone, luteinizing hormone releasing factor, alpha and beta seminal inhibins, and human growth hormone; genes that encode proteins such as enzymes, the absence of which leads to the occurrence of an abnormal state; genes encoding cytokines or lymphokines such as interferons, granulocytic macrophage colony stimulating factor, colony stimulating factor-1, tumor necrosis factor, and erythropoietin; genes encoding inhibitor substances such as alpha1-antitrypsin, genes encoding substances that function as drugs such as diphtheria and cholera toxins; and the like. Useful genes also include those useful for cancer therapies and to treat genetic disorders. Those skilled in the art have access to nucleic acid sequence information for virtually all known genes and can either obtain the nucleic acid molecule directly from a public depository, the institution that published the sequence, or employ routine methods to prepare the molecule.

[0038] For gene therapy use, the ligands described herein may be taken up in pharmaceutically acceptable carriers, such as, for example, solutions, suspensions, tablets, capsules, ointments, elixirs and injectable compositions. Pharmaceutical preparations may contain from 0.01 % to 99% by weight of the ligand. Preparations may be either in single or multiple dose forms. The amount of ligand in any particular pharmaceutical preparation will depend upon the effective dose, that is, the dose required to elicit the desired gene expression or suppression.

[0039] Suitable routes of administering the pharmaceutical preparations include oral, rectal, topical (including dermal, buccal and sublingual), vaginal, parenteral (including subcutaneous, intramuscular, intravenous, intradermal, intrathecal and epidural) and by naso-gastric tube. It will be understood by those skilled in the art that the preferred route of administration will depend upon the condition being treated and may vary with factors such as the condition of the recipient.

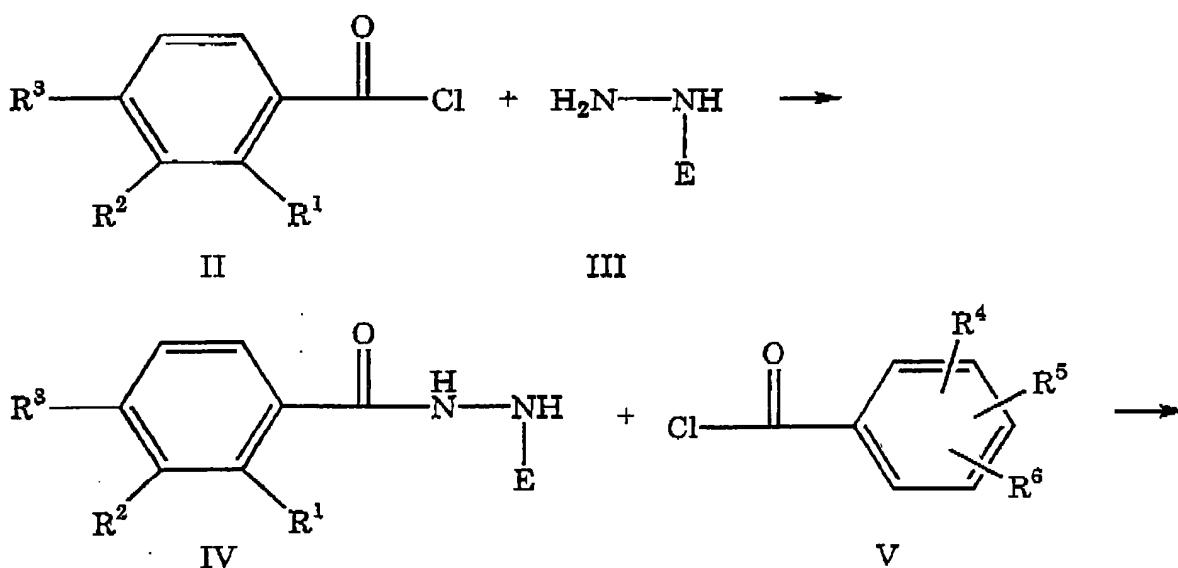
[0040] The ligands described herein may also be administered in conjunction with other pharmaceutically active compounds. It will be understood by those skilled in the art that pharmaceutically active compounds to be used in combination with the ligands described herein will be selected in order to avoid adverse effects on the recipient or undesirable interactions between the compounds. Examples of other pharmaceutically active compounds which may be used in combination with the ligands include, for example, AIDS chemotherapeutic agents, amino acid derivatives, analgesics, anesthetics, anorectal products, antacids and antifatulents, antibiotics, anticoagulants, antidotes, antifibrinolytic agents, antihistamines, anti-inflammatory agents, antineoplastics, antiparasitics, antiprotozoals, antipyretics, antiseptics, antispasmodics and anticholinergics, antivirals, appetite suppressants, arthritis medications, biological response modifiers, bone metabolism regulators, bowel evacuants, cardiovascular agents, central nervous system stimulants, cerebral metabolic enhancers, cerumenolytics, cholinesterase inhibitors, cold and cough preparations, colony stimulating factors, contraceptives, cytoprotective agents, dental preparations, deodorants, dermatologicals, detoxifying agents, diabetes agents, diagnostics, diarrhea medications, dopamine receptor agonists, electrolytes, enzymes and digestants, ergot preparations, fertility agents, fiber supplements, antifungal agents, galactorrhea inhibitors, gastric acid secretion inhibitors, gastrointestinal prokinetic agents, gonadotropin inhibitors, hair growth stimulants, hematinics, hemorrhheologic agents, hemostatics, histamine H₂ receptor antagonists, hormones, hyperglycemic agents, hypolipidemics, immunosuppressants, laxatives, leprostatics, leukapheresis adjuncts, lung surfactants, migraine preparations, mucolytics, muscle relaxant antagonists, muscle relaxants, narcotic antagonists, nasal sprays, nausea medications, nucleoside analogues, nutritional supplements, osteoporosis preparations, oxytocics, parasympatholytics, parasympathomimetics, Parkinsonism drugs, Penicillin adjuvants, phospholipids, platelet inhibitors, porphyria agents, prostaglandin analogues, prostaglandins, proton pump inhibitors, pruritus medications, psychotropics, quinolones, respiratory stimulants, saliva

stimulants, salt substitutes, sclerosing agents, skin wound preparations, smoking cessation aids, sulfonamides, sympatholytics, thrombolytics, Tourette's syndrome agents, tremor preparations, tuberculosis preparations, uricosuric agents, urinary tract agents, uterine contractants, uterine relaxants, vaginal preparations, vertigo agents, vitamin D analogs, vitamins, and medical imaging contrast media. In some cases the ligands may be useful as an adjunct to drug therapy, for example, to "turn off" a gene that produces an enzyme that metabolizes a particular drug.

[0041] For agricultural applications, in addition to the applications described above, the ligands described herein may also be used to control the expression of pesticidal proteins such as *Bacillus thuringiensis* (Bt) toxin. Such expression may be tissue or plant specific. In addition, particularly when control of plant pests is also needed, one or more pesticides may be combined with the ligands described herein, thereby providing additional advantages and effectiveness, including fewer total applications, than if the pesticides are applied separately. When mixtures with pesticides are employed, the relative proportions of each component in the composition will depend upon the relative efficacy and the desired application rate of each pesticide with respect to the crops, pests, and/or weeds to be treated. Those skilled in the art will recognize that mixtures of pesticides may provide advantages such as a broader spectrum of activity than one pesticide used alone. Examples of pesticides which can be combined in compositions with the ligands described herein include fungicides, herbicides, insecticides, miticides, and biocides.

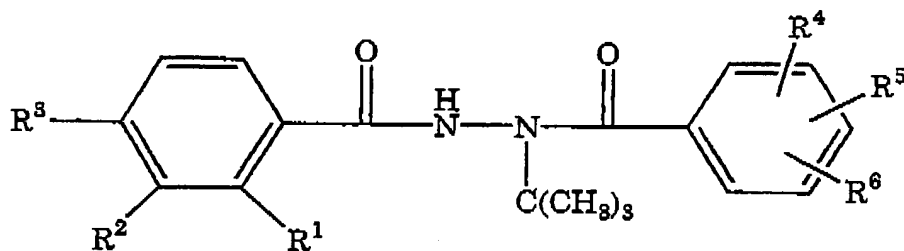
[0042] The ligands described herein can be applied to plant foliage as aqueous sprays by methods commonly employed, such as conventional high-liter hydraulic sprays, low-liter sprays, air-blast, and aerial sprays. The dilution and rate of application will depend upon the type of equipment employed, the method and frequency of application desired, and the ligand application rate. It may be desirable to include additional adjuvants in the spray tank. Such adjuvants include surfactants, dispersants, spreaders, stickers, antifoam agents, emulsifiers, and other similar materials described in McCutcheon's Emulsifiers and Detergents, McCutcheon's Emulsifiers and Detergents/Functional Materials, and McCutcheon's Functional Materials, all published annually by McCutcheon Division of MC Publishing Company (New Jersey). The ligands can also be mixed with fertilizers or fertilizing materials before their application. The ligands and solid fertilizing material can also be admixed in mixing or blending equipment, or they can be incorporated with fertilizers in granular formulations. Any relative proportion of fertilizer can be used which is suitable for the crops and weeds to be treated. The ligands described herein will commonly comprise from 5% to 50% of the fertilizing composition. These compositions provide fertilizing materials which promote the rapid growth of desired plants, and at the same time control gene expression.

[0043] The ligands described herein are known compounds or are easily prepared by one skilled in the art adapting the procedures described in U. S. Patents No. 5,117,057, 5,530,028, and 5,378,726. A two step process is typically used. In the first step, a substituted benzoic acid chloride of formula II is combined with a substituted hydrazine of formula III to give a monoacylhydrazine of formula IV. The monoacylhydrazine of formula IV is combined in a second step with another benzoic acid chloride of formula V resulting in formation of a compound of formula I.



[0044] The following examples demonstrate the activity of the ligands described herein.

[0045] The following ligands and comparative ligands were evaluated:



Ligand	R1	R2	R3	3-R4	5-R5	4-R6
CE-1	H	H	Me	Me	Me	
CE-2	H	H	H	H	H	
CE-3						
1	Et	OMe	H	OMe	H	
2	Me	-OCH ₂ CH ₂ O-		Me	Me	
3	Me	OMe	H	Me	Me	
4	Et	OMe	H	F	F	
5	Et	OMe	H	Me	Me	
6	Me	OEt	H	Me	Me	
7	Et	-OCH ₂ CH ₂ O-		Me	Me	
8	Et	-CH ₂ CH ₂ O-		Me	Me	
9	Me	OMe	H	Cl	Cl	
10	Et	OMe	H	Cl	Me	
11	i-Pr	OMe	H	Me	Me	
12	Et	OEt	H	Me	Me	
13	Et	OMe	H	Cl	Cl	
14	Me	OH	H	Me	Me	
15	Me	OH	H	Me	CH ₂ OH	
17	F	Et	Et	Me	Me	
18	Me	OMe	H	OMe	OMe	Me
CE-1= tebufenozide						
CE-2 = 1,2-dibenzoyl-1- <i>tert</i> -butyl- hydrazine						
CE-3 = muristerone A						

Gene Constructs

[0046] pVgRXR (Invitrogen Corp., Carlsbad, California) is an 8728 kb plasmid that expresses both VgEcR and RXR to form a modified heterodimeric nuclear receptor (see No, D., et. al., (1996) Proc. Natl. Acad. Sci. USA, 93, 3346-3351). The ecdysone receptor (VgEcR) is derived from the natural *Drosophila* ecdysone receptor and modified to contain the VP16 transactivation domain (see Cress, W.D., and Triezenberg, S.J. (1991), Science 251, 87-90; Sadowski, I., et. al (1988) Nature 335, 563-564; Triezenberg, S.J. et. al., (1988), Genes Dev 2, 718-729; Triezenberg, S.J., et. al., (1988),

Genes Dev 2, 730-742). RXR, is the mammalian homologue of USP (ultraspiracle), the natural partner to the *Drosophila* ecdysone receptor (see Yao, T.-P., et. al., (1993), Nature 366, 476-479; Yao, T.-P., et. al., (1992), Cell 71, 63-72). The P-box region of the VgEcR DNA binding domain was modified to recognize the hybrid ecdysone response element that consists of one half-site from the glucocorticoid response element (see Umesono, K., and Evans, R. M. (1989), Cell 57, 1139-1146) and one half-site from the natural ecdysone response element. This hybrid response element reduces any possible interaction with the farnesoid X receptor that may bind the natural ecdysone responsive element (EcRE). The cytomegalovirus enhancer-promoter drives expression of VgEcR, and the Rous sarcoma virus promoter drives expression of RXR. The vector pIND (Invitrogen Corp.) is a 5024 bp vector based on pcDNA3.1. It contains five hybrid E/GREs recognized by the modified ecdysone receptor expressed from pVgRXR, and a minimal heat shock promoter (see Yao, T. P., et. al., (1993), Nature, 366, 476-479). pIND/lacZ (Invitrogen Corp.) is a 8170 bp plasmid that contains the β -galactosidase gene as a reporter enzyme. Cotransfection of pIND/lacZ and pVgRXR, results in the induction of β -galactosidase expression upon addition of ecdysone agonists such as muristerone A. The reporter plasmid pIND/luc was constructed by subcloning the firefly luciferase gene from pGL3 (Promega/E1741) as a Nhe I - BamHI fragment into pIND, also digested with Nhe I and BamHI.

Maintenance of Mammalian Cell Lines and Transfection

[0047] CHO cells (ATCC # CCL-61) were maintained in F-12 Nutrient mixture (Ham's media) (Gibco/BRL, 11765-054) supplemented with 10% fetal bovine serum (FBS, complete medium; Gibco/BRL, 16000-036). These cells were maintained at 37°C in 5% CO₂ in 95% atmospheric air.

[0048] The CHO cells were seeded into 12-well tissue culture plates at a concentration of 0.5 X 10⁵ cells per ml per well. The lipid Pfx-8 (Invitrogen, T930-18) was used to transiently transfect the ecdysone inducible expression system (Invitrogen catalog number K1000-01). The ecdysone inducible expression system consists of pVgRXR, encoding the receptor subunits, and pIND/lacZ or pIND/luc, containing the response element and a reporter gene encoding β -galactosidase or luciferase respectively. Twenty-four hours after seeding cells, the lipid/DNA solution was prepared by 0.5 mg/ml of VgRXR, and 0.5 mg/ml pIND/lacZ or pIND/luc DNAs in sterile water. Thirty-six μ l of Pfx-8 was diluted in a sterile 17 x 100 mm polystyrene tube containing 1.5 ml Opti-MEM medium (Gibco/BRL, 31985). Six μ l of the plasmid DNA stock was mixed with 1.5 ml of Opti-MEM medium in another polystyrene tube. The DNA and the lipid solutions were combined to make 3 ml of transfection solution (enough for one set of triplicate samples). Cells were washed with phosphate buffered solution (PBS; Gibco/BRL, 14190-144) three times by aspirating medium from the cells. One ml transfection solution was added per well. Cells were incubated for four hours, and the transfection medium replaced with the same volume of complete medium. Cells were incubated for an additional 20 hours.

Stably Transformed Mammalian Cell Line

[0049] CHO cells stably transformed with pVgRXR and pIND/LacZ(SPI) (Invitrogen Corp.) were maintained in Hams F-12 media containing 10% FBS, 2 mM glutamine, 300 μ g/ml Zeocin (Invitrogen Corp.) and 300 μ g/ml Hygromycin B. pIND(SPI), an alternative to the pIND vector, can be induced to absolute expression levels five-fold greater than that of pIND due to the presence of 3 cis-acting SPI elements. Basal expression levels are correspondingly greater.

Treatment with Ligands

[0050] Stock solutions (10⁻² M) were prepared for muristerone A and the non-steroidal ligands described herein in ethanol and acetone respectively, and stored at -20°C. Twenty-four hours after transfection, a test compound was added at a final concentration of 10 μ M (10⁻⁵ M) to each 1 ml cell culture well. Muristerone A (Sigma, M7888), at 10 μ M concentration was used as a positive control. Acetone alone was added as a negative control.

Reporter Gene Assays

[0051] Reporter gene expression was evaluated 24-48 hours after treatment of transiently transfected cells or 24 hours after treatment of the stably transformed cell line with test ligands. β -galactosidase was assayed by either staining fixed cells or assaying for enzymatic activity in cellular lysates. β -galactosidase catalyzes the hydrolysis of the β -galactoside, X-gal (5-bromo-4-chloro-3-indolyl- β -galactopyranoside) producing a blue color within the fixed cells (Invitrogen, K1465-01) that can be visualized under a microscope. Alternatively, reporter lysis buffer (RLB; Promega/E397A) was used to lyse the cells for the detection of β -galactosidase activity using a chemiluminescent substrate, Galacto-Star (Tropix/BM100S). The cells within each 12-well plate were lysed with 250 μ l of RLB. Twenty μ l of each extract was assayed with 100 μ l of substrate.

[0052] For detection of luciferase activity, cells were washed twice with PBS and lysed with 250 μ l RLB per well. After

10 minutes, the plate was frozen at -80°C for 10 minutes and returned to room temperature. A 20 µl sample of lysate was mixed with 100 µl luciferase assay reagent (Promega, E1500). For both β-galactosidase and luciferase assays, luminescence was detected at room temperature using a DYNEX MLX microtiter plate luminometer equipped with an autoinjector for the delivery of the substrate.

Cytosolic ecdysone receptor extract preparation from *Drosophila* Kc cells

[0053] The dipteran cell line Kc167, originally derived from *Drosophila* embryos (see Echalié, G. and Ohanessian, A. (1969) C. R. Acad. Sci., 268, 1771) was obtained from Dr. Peter Cherbas (Indiana University) and maintained as described (Cherbas, L., et. al., (1994), Methods in Cell Biology, 44, 161-179). A 400 ml culture of Kc cells (3 x 10⁷ cells per ml) was centrifuged at 700 x g for 10 minutes at room temperature to pellet the cells. The supernatant was aspirated and the pellet was resuspended in 70 ml of cold TM buffer (10mM Tris, 5 mM MgCl₂, 1mM DTT, pH 7.2). After a 10 minute incubation on ice, the cells were centrifuged at 2,300 x g. The supernatant was discarded. The cell pellet was frozen at -20°C for 1 hour. The frozen cell pellet was thawed slowly on ice and homogenized in a cold Potter-Elvehjem homogenizer using a teflon pestle on a Caframo homogenizer motor, at setting 500 with 10 up and down strokes at 4°C. This slurry was centrifuged at 100,000 x g in a swinging bucket rotor for 60 minutes. The supernatant, containing the cytosolic protein extract was diluted with T buffer (10mM Tris, 1 mM DTT, pH 7.2) to a protein concentration of 5 mg / ml. This extract was used immediately for ligand binding assays.

Nuclear ecdysone receptor preparation from *Plodia interpunctella* cells

[0054] The cell line IAL-PID2, derived from imaginal wing discs of the lepidopteran *Plodia interpunctella* was obtained from H. Oberlander and maintained as described (Lynn, E. E. and Oberlander, H., (1983), J. Insect Physiology, 29, 591-96). A 300 ml stationary phase culture of *P. interpunctella* cells was centrifuged at 700 x g for 10 min at room temperature to pellet the cells. The supernatant was aspirated and the cellular pellet resuspended in 35 ml TMT (TM buffer with 0.1% Triton X-100). The suspension was homogenized with twenty up and down strokes of a Dounce homogenizer on ice. Homogenate was incubated on ice for 10 minutes and then centrifuged for 15 min at 900 x g. The pellet was resuspended in 15 ml TM buffer and centrifuged at 2,300 x g. This pellet was extracted in TMK buffer (TM buffer with 800 mM KCl) by using a glass rod to crush the pellet until a gelatinous slurry was formed and incubated on ice for 15 minutes. The extract was centrifuged at 100,000 x g in a swinging bucket rotor for 60 minutes. The supernatant, which constituted the nuclear extract, was desalted on a 10 DG desalting column (Bio-Rad, 732-2010) equilibrated with T-buffer. The total protein concentration in the nuclear extract was adjusted to 5 mg per ml with T-buffer containing 1 mM DTT.

Bacterial Glutathion-S-transferase Fusion Protein Preparation

[0055] Bacterial glutathion-S-transferase (GST) fusion proteins of the spruce budworm (*Choristoneura fumiferana*) EcR (CfEcR), containing CDEF domains only, and ultraspiracle protein (CfUSP) were also used in radioligand displacement assays. cDNA encoding the CDEF domains of CfEcR were constructed following PCR amplification using primers that contain BamHI and EcoRI sites (Perera SC, M Sundaram, Krell PJ, Retnakaran A, Dhadialla TS, SR Palli, 1999 Arch. Insect Biochem. Physiol. 41, In Press). The PCR product was digested with BamHI and EcoRI and cloned into pGEX-3X vector obtained from Pharmacia Biotech. CfUSP coding region was amplified using primers that contain BamHI and EcoRI sites and cloned into pGEX-2T vector obtained from Pharmacia Biotech. *E. coli* transformed with each of the two vectors were grown and induced to produce the fusion proteins as detailed in the Pharmacia Biotech GST- technical bulletin.

Ecdysone receptor competitive binding assay

[0056] ³H ponasterone A, a potent phytoecdysteroid, (66,000 dpm; specific activity 170 Ci/mmol; NEN Life Science Products, Boston, Massachusetts) was mixed with 100 µl Kc cytosolic or *Plodia* nuclear ecdysone receptor extracts in 0.8 X 50 mm glass test tubes in the absence or presence of 10 µM unlabelled 20-hydroxyecdysone (20E) to obtain an estimate of total or non-specific, respectively, binding of tritiated ponasterone A. The tubes were vortexed and incubated overnight at 4°C for Kc cytosolic extracts or 1.5 hours for *Plodia* nuclear extracts for binding reactions to reach equilibrium. At the end of equilibrium binding times, the bound tritiated ponasterone A was separated from the unbound by addition of ice-cold 600 µl or 300 µl dextran-coated charcoal solution (500 mg of Sigma HCL-washed activated charcoal, 50 mg Pharmacia Dextran T70, 50 ml T-buffer) to Kc or *Plodia* extracts, respectively. The tubes were vortexed briefly and centrifuged at 7,000 x g to pellet the charcoal. The supernatant, 600 µl or 300 µl for Kc or *Plodia* extract reactions, respectively, containing tritiated ponasterone A bound to proteins, was aspirated into liquid scintillation vials containing

5 ml scintillation cocktail (ReadySafe®, Beckmann) each. The mixture was vortexed and the amount of total or non-specifically bound radioactivity measured in a Beckman LS500 liquid scintillation counter with a 60% counting efficiency for tritium.

Determination of K_d values for competitive inhibitors of ³H-ponasterone A binding to ecdysteroid receptor complexes in Kc or *Plodia* cell extracts and bacterial fusion proteins

[0057] The concentrations of competitors that inhibited 50% of tritiated ponasterone A binding (IC₅₀) was determined by incubating Kc or *Plodia* nuclear extracts or extracts of bacteria producing CfEcR(CDEF)-GST and CfUSP-GST fusions proteins with ³H-ponasterone A (66,000 dpm per reaction) in the presence of a range of concentrations (0.1 nM to 10 μM) of test compounds. In the case of binding reactions using bacterial fusion proteins, 20 or one μl extract of bacteria producing CfEcR(CDEF)-GST or CfUSP-GST fusion proteins, respectively, were included per binding reaction. The assay conditions and determination of total and non-specific binding was as described above. The volume of competitor in solvent, like that for 20E or solvent alone, was kept at 1% of total reaction volume by using a 100 fold concentrated stock solution (i.e. a 100 fold dilution of stock solutions). The remainder of the assay was as described above. Each reaction was carried out in duplicate per concentration per test compound. Specific binding of ³H-ponasterone A was determined by subtracting non-specific binding (that obtained in the presence of 10 μM 20E) from total (no competitor) or competed radioactivity bound. The data for each test compound was analyzed using IGOR Pro software (WaveMetrics, Lake Oswego, OR) to calculate IC₅₀ values. The binding constant (K_d in μM) for the test compounds was calculated by incorporating the Cheng-Prusoff equation (see Munson P.J. and Rodbard D. (1980) Anal. Biochem. 107, 220-239) in the IGOR Pro software.

[0058] Tables 1- 3 summarize the data obtained:

Table 1

Ligand	Kc log(1/EC50)	Plodia log(1/IC50)	Bacterial CfECR log(1/IC50)
CE-1	6.55	8.7	
CE-2	5.52	6.48	
CE-3	7.96		
1	7.36	8.52	
2	7.16	9.14	
3	6.73	9.04	8.76
4	7.41	8.51	
5	7.70	9.49	
6		8.82	
7		8.78	
8		8.85	
9		9.00	
10		8.80	
11		8.93	
12	8.10	9.63	
13	7.44	9.06	
14			8.61
15			8.49
17		9.24	
18		8.82	

Table 2

Luciferase Gene Activation in Transiently Transfected CHO Cells			
Ligand 10 μ M	Luciferase activity ¹	Induction Ratio ²	Kd for Kc cells (nM)
none	42		
CE-1	89	2	110
CE-2	36	1	2000
CE-3	316	8	2.3
CE-4	418	10	0.7
1	275	7	39.5
2	204	5	60
3	324	8	124
4	251	6	47
5	313	8	13.3
12	305	7	5.3
13	226	5	65
CE-4 = ponasterone A ¹ In relative light units (RLU) - Average of duplicate samples ² Ratio of RLU in presence and absence of the ligand			

Table 3

β -Galactosidase Gene Activation in Stably Transformed CHO Cells			
Ligand 10 mM	β -Galactosidase activity ¹	Induction Ratio ²	Kd for Kc cells (nM)
none	200		
CE-1	323	2	192
CE-2	145	1	2000
CE-3	9830	49	2
CE-4	10386	52	0.7
1	1123	6	40
2	536	3	47
3	681	3	124
4	1963	10	26
5	11599	58	13
12	9830	49	5
13	6667	33	24
¹ In relative light units (RLU) - Average of triplicate samples ² Ratio of RLU in presence and absence of the ligand			

[0059] These data indicate that the ligands described herein are able to induce gene expression at concentrations comparable to or less than those of the known compounds CE-1 and CE-2.

[0060] In addition to improved gene expression modulation capabilities, the ligands are expected to have greater utility in intact plants and animals due to their superior transport and metabolic properties. The following example demonstrates the improved plant transport of a ligand described herein (Ligand 3) compared to a known ligand (CE-1) in plants.

Translaminar Movement

[0061] A study was conducted to evaluate translaminar movement of two compounds into cotton leaves using neonate beet armyworm larvae, *Spodoptera exigua*, as the model system.

[0062] Two treatments were evaluated: emulsifiable concentrates of CE-1 and Ligand 3 (5% and 19%, respectively), at a concentration of 100 µg/ml to simulate spray tank concentrations. These treatments were then compared with plants treated with a chemical standard known to have excellent translaminar movement (emamectin benzoate, 0.16% EC [20 µg/ml], Merck & Co.) and with nontreated plants.

[0063] Four-week-old cotton, *Gossypium hirsutum* L. cv. Stoneville, plants were treated with each treatment by painting the top surface of a single replicate leaf/plant with a corresponding treatment. Plants were held in a controlled-environment glasshouse maintained at 27 °C until needed. Residual effectiveness was evaluated by challenging treated and nontreated foliage with neonate beet armyworm larvae, *Spodoptera exigua* (Hübner), on 3 dates: 1, 7, and 14 days after treatment (DAT). On each sample date, the single treated leaf/plant was excised from each of five replicate plants per treatment. Leaves were then fastened to the top lid of a plastic Petri dish (100 x 20 mm) and then infested with 10 first instar larvae. First instar beet armyworm larvae commonly feed on the under surface of leaves and generally do not feed all the way through to the upper surface. Thus translaminar movement of material would be necessary to affect the larvae since the compounds were applied to the leaf top-surface only. Mortality of larvae was recorded 4 days after infestation. In addition, any live larvae were observed for symptoms characteristic of molt accelerating compounds (formation of new cuticle and/or slipped head capsule). Each treatment was replicated five times.

[0064] Percentage mortality data were transformed to the arcsine of the square-root and then analyzed by ANOVA using JMP (Ver. 3.2.1). Means were separated by the Tukey-Kramer test (P = 0.05).

[0065] Results showed that Ligand 3 was significantly more effective than CE-1 at killing *S. exigua* larvae from translaminar movement and, therefore, would be expected to have higher systemicity in plants.

Table 4
Percentage Mortality

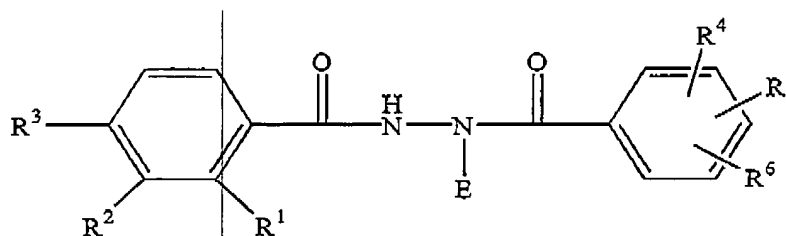
Treatment	Conc. (µg/ml)	Day 1	After 7	Applic. 14
CE-1, 5% EC	100	2.5	0.0	22.7
3, 19% EC	100	100.0	93.8	100.0
Emamectin benzoate 0.16 EC + AG-98	20	100.0	100.0	100.0
Nontreated check		11.6	6.5	8.1

AG-98 = 0.12% of the nonionic surfactant Latron AG-98 surfactant (Rohm and Haas Company)

Claims

1. An *in vitro* method to modulate exogenous gene expression comprising contacting an ecdysone receptor complex comprising:

- a) a DNA binding domain;
- b) a ligand binding domain;
- c) a transactivation domain; and
- d) a ligand of the formula:



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt;

provided that:

a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl;

c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is:

i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or

ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, CN, OMe, or OEt;

d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;

then R¹ is Me or Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

with a DNA construct comprising:

a) the exogenous gene; and

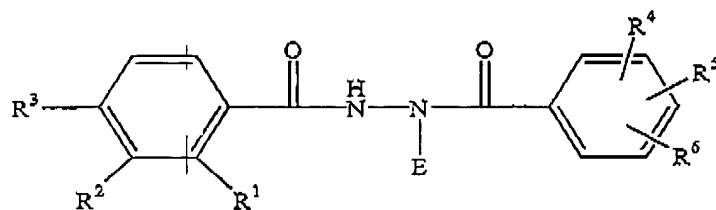
b) a response element;

wherein:

a) the exogenous gene is under the control of the response element; and

b) binding of the DNA binding domain to the response element in the presence of the ligand results in activation or suppression of the gene.

2. A method to modulate the expression of one or more exogenous genes in a plant, comprising administering to the plant an effective amount of a ligand of the formula:



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂O CN, OMe, or OEt;

provided that:

a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl;

c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is:

i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-1-5-Me, 2-Cl, or 3-Me; or

ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, CN, OMe, or OEt;

d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

h) when R² is Me, Et n-Pr, i-Pr, Ac, F, Cl, OH, or SMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

wherein the cells of the plant contain:

a) an ecdysone receptor complex comprising:

1) a DNA domain;

2) a binding domain for the ligand; and

3) a transactivation domain; and

b) a DNA construct comprising:

1) the exogenous gene; and

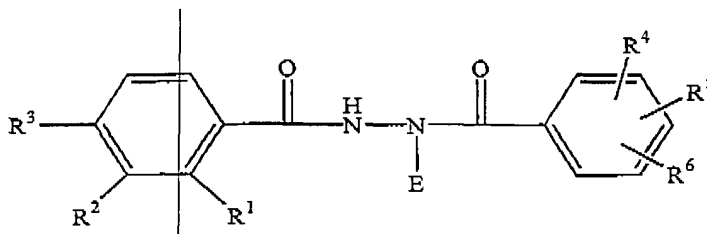
2) a response element; and

wherein:

- a) the exogenous gene is under the control of the response element; and
- b) binding of the DNA binding domain to the response element in the presence of the ligand results in activation or suppression of the gene.

3. A method for producing a polypeptide comprising the steps of:

- a) selecting a cell which is insensitive to exposure to a ligand of the formula:



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt;

provided that:

- a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

- b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl;

- c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is:

- i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or

- ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, OMe, or OEt;

- d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

- e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

- f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring; then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

- g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

- h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

b) introducing into the cell:

1) a DNA construct comprising:

- a) an exogenous gene encoding the polypeptide; and
b) a response element;

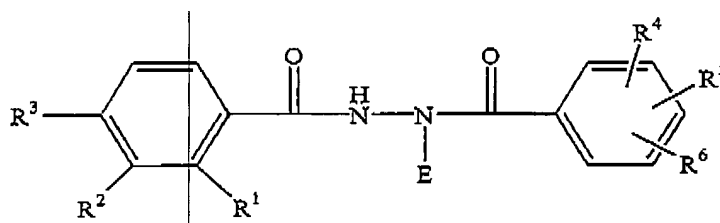
2) an ecdysone receptor complex comprising:

- a) a DNA binding domain;
b) a binding domain for the ligand; and
c) a trans activation domain; and

wherein the gene is under the control of the response element; and

c) exposing the cell to the ligand of step(a).

4. A method for regulating endogenous or heterologous gene expression in a transgenic plant comprising contacting a ligand of the formula:



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently, H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt;

provided that:

a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl ;

c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is:

i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or

ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, CN, OMe, or OEt;

d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

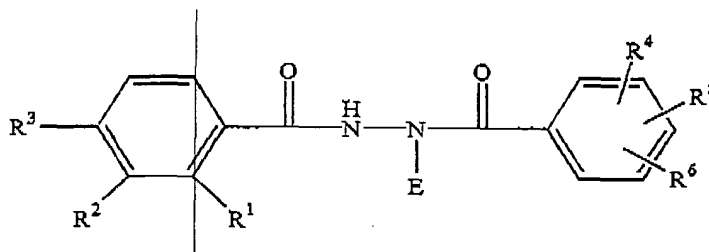
then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SOMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

with an ecdysone receptor complex within the cells of the plant, wherein the cells further contain a DNA binding sequence for the ecdysone receptor complex when in combination with the ligand and wherein formation of an ecdysone receptor complex-ligand-DNA binding sequence complex induces expression of the gene.

5. The method of any one of claims 1 to 4 wherein the ligand is of the specified formula and E is t-butyl.
6. The method of any one of claims 1 to 5 wherein the ligand is of the specified formula and R¹ is Me, Et, i-Pr or F.
7. The method of any one of claims 1 to 6 wherein the ligand is of the specified formula and R² is OH, OMe, OEt, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy or dihydrofuryl ring with the oxygen adjacent to a phenyl carbon.
8. The method of any one of claims 1 to 7 wherein the ligand is of the specified formula and R³ is H, Et or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy or dihydrofuryl ring with the oxygen adjacent to a phenyl carbon.
9. The method of any one of claims 1 to 4 wherein the ligand is of the specified formula and E is t-butyl, R¹ is Et, R² is OEt, R³ is H, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me.
10. The method of any one of claims 1 to 9 wherein the ecdysone receptor complex is a chimeric ecdysone receptor complex and the DNA construct further comprises a promoter.
11. Use of a ligand of the formula (I):



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt; provided that:

a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl;

c) when R¹ is Et and R² is OMe or OEt;
then R³ is H and the combination R⁴, R⁵, and R⁶ is:

- i) 3,5-di- OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or
ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH CN, OMe, or OEt;

d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;
then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SOMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

for modulating *in vitro* the expression of one or more exogenous genes in an animal cell, wherein the cell contains:

a) an ecdysone receptor complex comprising:

- 1) a DNA binding domain;
2) a binding domain for the ligand; and
3) a transactivation domain; and

b) a DNA construct comprising:

- 1) the exogenous gene; and
2) a response element; and

wherein:

- a) the exogenous gene is under the control of the response element; and
b) binding of the DNA binding domain to the response element in the presence of the ligand results in activation or suppression of the gene.

12. Use according to claim 11 wherein the ligand is of the specified formula and E is t-butyl.

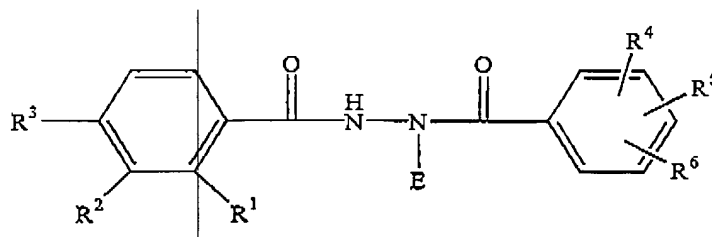
13. Use according to claim 11 or 12 wherein the ligand is of the specified formula and R¹ is Me, Et, i-Pr or F.

14. Use according to any one of claims 11 to 13 wherein the ligand is of the specified formula and R² is OH, OMe, OEt, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy or dihydrofuryl ring with the oxygen adjacent to a phenyl carbon.

15. Use according to any one of claims 11 to 14 wherein the ligand is of the specified formula and R³ is H, Et or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy or dihydrofuryl ring with the oxygen adjacent to a phenyl carbon.

16. Use according to any one of claims 11 to 15 wherein the ligand is of the specified formula and E is t-butyl, R¹ is Et, R² is OEt, R³ is H, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me.

17. Use of a ligand of the formula (I):



wherein:

E is a (C₄-C₆)alkyl containing a tertiary carbon;

R¹ is H, Me, Et, i-Pr, F, or OMe;

R² is H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SMe, or joined with R³ and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R³ is H, Et, or joined with R² and the phenyl carbons to which R² and R³ are attached to form an ethylenedioxy, a dihydrofuryl ring with the oxygen adjacent to a phenyl carbon, or a dihydropyryl ring with the oxygen adjacent to a phenyl carbon;

R⁴, R⁵, and R⁶ are independently H, Me, Et, F, Cl, CH₂OH, CN, OMe, or OEt;

provided that:

a) when R¹ is Me and R² is OMe;

then R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, or 3,5-di-F;

b) when R¹ is Me and R² is OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl ;

c) when R¹ is Et and R² is OMe or OEt;

then R³ is H and the combination R⁴, R⁵, and R⁶ is:

i) 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- or 2,5-di-F, 2,4- or 2,5-di-Cl, 3,OMe, 2-Cl-5-Me, 2-Cl, or 3-Me; or

ii) R⁶ is H, R⁴ is Me, and R⁵ is Et, F, Cl, CH₂OH, CN, OMe, or OEt;

d) when R¹ is i-Pr;

then R² is OMe, or OEt; R³ is H; and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

e) when R³ is Et;

then R² is H, R¹ is F, and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

f) when R² and R³, together with the phenyl carbons to which they are attached, form an ethylenedioxy ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me;

g) when R² and R³, together with the phenyl carbons to which they are attached, form a dihydrofuryl or dihydropyryl ring;

then R¹ is Et and the combination R⁴, R⁵, and R⁶ is 3,5-di-Me; and

h) when R² is Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, or SMe;

then R¹ is Et, R³ is H, the combination R⁴, R⁵, and R⁶ is 3,5-dx-IIQe;

in the preparation of pharmaceutical for modulating the expression of one or more exogenous genes in an animal, wherein cells of the animal contain:

a) an ecdysone receptor complex comprising:

1) a DNA binding domain;

2) a binding domain for the ligand; and

3) a transactivation domain; and

b) a DNA construct comprising:

- 1) the exogenous gene; and
- 2) a response element; and

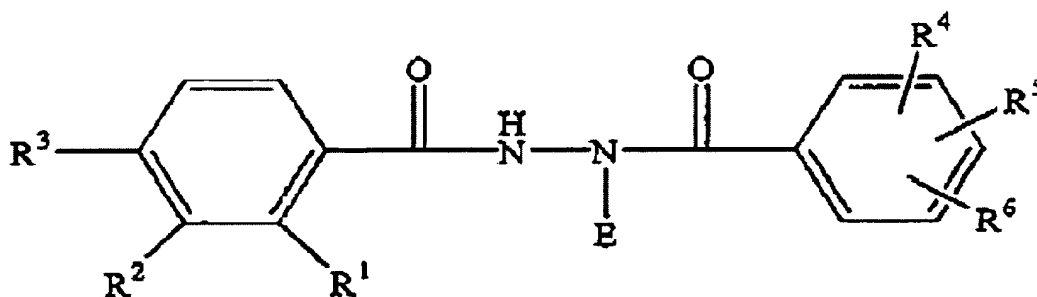
wherein:

- a) the exogenous gene is under the control of the response element; and
- b) binding of the binding domain to the response element in the presence of the ligand results in activation or suppression of the gene.

Patentansprüche

1. *In vitro*-Verfahren zum Modulieren von exogener Genexpression, umfassend das Inkontaktbringen eines Ecdyson-Rezeptorkomplexes, welcher folgendes umfaßt:

- a) eine DNA-Bindungsdomäne,
- b) eine Ligandenbindungsdomäne,
- c) eine Transaktivierungsdomäne und
- d) einen Liganden der Formel



wobei:

E ein (C₄-C₆)-Alkyl ist, welches einen tertiären Kohlenstoff enthält,
 R¹ H, Me, Et, i-Pr, F oder OMe ist,
 R² H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe ist oder verbunden ist mit R³ und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder einen Dihydropyrylring, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,
 R³ H, Et ist oder verbunden ist mit R² und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylring, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,
 R⁴, R⁵ und R⁶ unabhängig voneinander H, Me, Et, F, Cl, CH₂OH, CN, OMe oder OEt sind, mit der Maßgabe, daß

- a) wenn R¹ Me ist und R² OMe ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl oder 3,5-Di-F ist,
- b) wenn R¹ Me ist und R² OEt ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-D-Cl ist,
- c) wenn R¹ Et ist und R² OMe oder OEt ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ folgendes ist:

- i) 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl, 3-OMe, 2-Cl-5-

Me, 2-Cl oder 3-Me oder

ii) R⁶ H ist, R⁴ Me ist und R⁵ Et, F, Cl, CH₂OH, CN, OMe oder OEt ist,

d) wenn R¹ i-Pr ist,

dann R² OMe oder OEt ist, R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

e) wenn R³ Et ist,

dann R² H ist, R¹ F ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

f) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Ethylendioxyring bilden,

dann R¹ Me oder Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

g) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Dihydrofuryl- oder Dihydropyrylring bilden,

dann R¹ Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist, und

h) wenn R² Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH oder SOMe ist,

dann R¹ Et ist, R³ H ist, die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist, mit einem DNA-Konstrukt, welches folgendes umfaßt:

a) das exogene Gen und

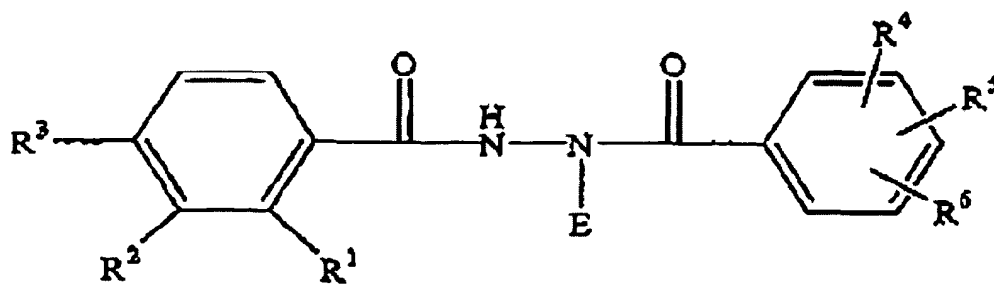
b) ein Antwortelement,

wobei:

a) das exogene Gen der Kontrolle durch das Antwortelement untersteht und

b) eine Bindung der DNA-Bindungsdomäne an das Antwortelement in der Gegenwart des Liganden zu einer Aktivierung oder Unterdrückung des Gens führt.

2. Verfahren zum Modulieren der Expression eines oder mehrerer exogener Gene in einer Pflanze, welches das Verabreichen einer wirksamen Menge eines Liganden der nachfolgenden Formel an die Pflanze umfaßt:



wobei:

E ein (C₄-C₆)-Alkyl ist, welches einen tertiären Kohlenstoff enthält,

R¹ H, Me, Et, i-Pr, F oder OMe ist,

R² H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe ist oder verbunden ist mit R³ und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,

R³ H, Et ist oder verbunden ist mit R² und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, R⁴, R⁵ und R⁶ unabhängig voneinander H, Me, Et, F, Cl, CH₂OH, CN, OMe oder OEt sind,

mit der Maßgabe, daß:

a) wenn R¹ Me ist und R² OMe ist,

dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl oder 3,5-Di-F ist,

b) wenn R^1 Me ist und R^2 OEt ist,
dann R^3 H ist und die Kombination aus R^4 , R^5 und R^6 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4-
oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl ist,
c) wenn R^1 Et ist und R^2 OMe oder OEt ist,
dann R^3 H ist und die Kombination aus R^4 , R^5 und R^6 folgendes ist:

- i) 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl
oder 3-Me oder
- ii) R^6 H ist, R^4 Me ist und R^5 Et, F, Cl, CH_2OH , CN, OMe oder OEt ist,

d) wenn R^1 i-Pr ist,
dann R^2 OMe oder OEt ist, R^3 H ist und die Kombination aus R^4 , R^5 und R^6 3,5-Di-Me ist,
e) wenn R^3 Et ist,
dann R^2 H ist, R^1 F ist und die Kombination aus R^4 , R^5 und R^6 3,5-Di-Me ist,
f) wenn R^2 und R^3 zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Ethylendioxyring
bilden,
dann R^1 Et ist und die Kombination aus R^4 , R^5 und R^6 3,5-Di-Me ist,
g) wenn R^2 und R^3 zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Dihydrofuryl-
oder Dihydropyrylring bilden,
dann R^1 Et ist und die Kombination aus R^4 , R^5 und R^6 3,5-Di-Me ist, und
h) wenn R^2 Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH oder SOMe ist,
dann R^1 Et ist, R^3 H ist und die Kombination aus R^4 , R^5 und R^6 3,5-Di-Me ist,

wobei die Zellen der Pflanze folgendes enthalten:

a) einen Ecdyson-Rezeptorkomplex, welcher folgendes umfaßt:

- 1 eine DNA-Bindungsdomäne,
- 2 eine Bindungsdomäne für den Liganden und
- 3 eine Transaktivierungsdomäne und

b) ein DNA-Konstrukt, welches folgendes umfaßt:

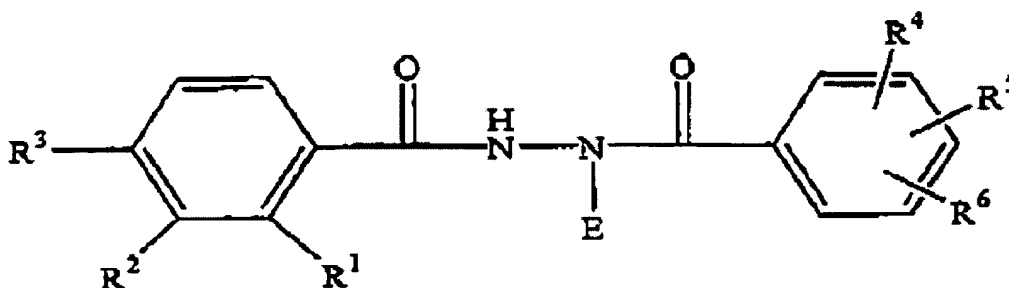
- 1) das exogene Gen und
- 2) ein Antwortelement, und

wobei:

- a) das exogene Gen der Kontrolle durch das Antwortelement untersteht und
- b) eine Bindung der DNA-Bindungsdomäne an das Antwortelement in der Gegenwart des Liganden zu einer
Aktivierung oder Unterdrückung des Gens führt.

3. Verfahren zur Herstellung eines Polypeptids, wobei das Verfahren die folgenden Stufen umfaßt:

a) Auswählen einer Zelle, die gegenüber Aussetzung an einen Liganden der folgenden Formel unempfindlich ist:



wobei:

E ein (C₄-C₆)-Alkyl ist, welches einen tertiären Kohlenstoff enthält,

R¹ H, Me, Et, i-Pr, F oder OMe ist,

R² H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, S_{OMe} ist oder verbunden ist mit R³ und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,

R³ H, Et ist oder verbunden ist mit R² und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,

R⁴, R⁵ und R⁶ unabhängig voneinander H, Me, Et, F, Cl, CH₂OH, CN, OMe oder OEt sind,

mit der Maßgabe, daß:

a) wenn R¹ Me ist und R² OMe ist,

dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl oder 3,5-Di-F ist,

b) wenn R¹ Me ist und R² OEt ist,

dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl ist,

c) wenn R¹ Et ist und R² OMe oder OEt ist,

dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ folgendes ist:

i) 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl oder 3-Me oder

ii) R⁶ H ist, R⁴ Me ist und R⁵ Et, F, Cl, CH₂OH, OMe oder OEt ist,

d) wenn R¹ i-Pr ist,

dann R² OMe oder OEt ist, R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

e) wenn R³ Et ist,

dann R² H ist, R¹ F ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

f) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Ethylendioxyring bilden,

dann R¹ Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

g) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Dihydrofuryl- oder Dihydropyrylring bilden,

dann R¹ Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist, und

h) wenn R² Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH oder S_{OMe} ist,

dann R¹ Et ist, R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

b) Einbringen der folgenden in die Zelle:

1) ein DNA-Konstrukt, welches folgendes umfaßt:

a) ein exogenes Gen, welches das Polypeptid codiert, und

b) ein Antwortelement,

wobei das Gen der Kontrolle durch das Antwortelement unterliegt, und

2) einen Ecdyson-Rezeptorkomplex, welcher folgendes umfaßt:

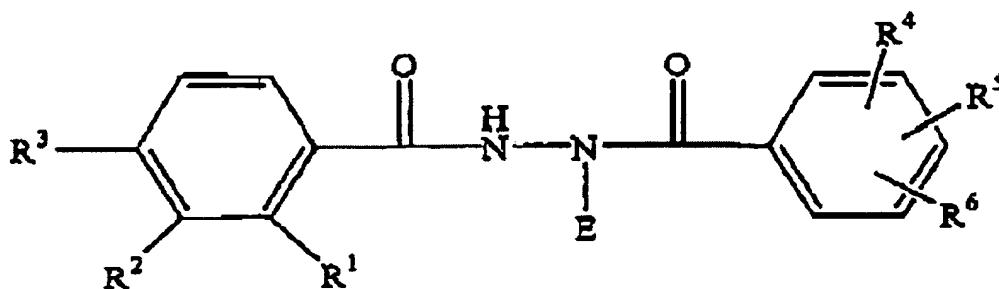
a) eine DNA-Bindungsdomäne,

b) eine Bindungsdomäne für den Liganden und

c) eine Transaktivierungsdomäne, und

c) Aussetzen der Zelle an den Liganden aus Stufe (a).

4. Verfahren zum Regulieren der endogenen oder heterologen Genexpression in einer transgenen Pflanze, umfassend das Inkontaktbringen eines Liganden der Formel:



wobei:

E ein (C₄-C₆)-Alkyl ist, welches einen tertiären Kohlenstoff enthält,
 R¹ H, Me, Et, i-Pr, F oder OMe ist,
 R² H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe ist oder verbunden ist mit R³ und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,
 R³ H, Et ist oder verbunden ist mit R² und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,
 R⁴, R⁵ und R⁶ unabhängig voneinander H, Me, Et, F, Cl, CH₂OH, CN, OMe oder OEt sind,

mit der Maßgabe, daß:

a) wenn R¹ Me ist und R² OMe ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl oder 3,5-Di-F ist,
 b) wenn R¹ Me ist und R² OEt ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl ist,
 c) wenn R¹ Et ist und R² OMe oder OEt ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ folgendes ist:

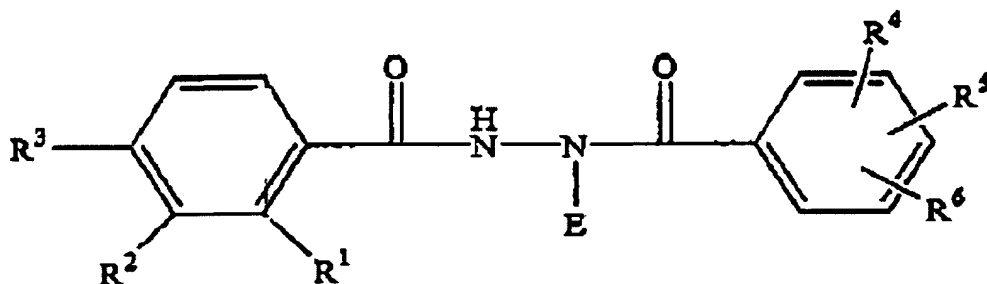
i) 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl oder 3-Me oder
 ii) R⁶ H ist, R⁴ Me ist und R⁵ Et, F, Cl, CH₂OH, CN, OMe oder OEt ist,

d) wenn R¹ i-Pr ist,
 dann R² OMe oder OEt ist, R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,
 e) wenn R³ Et ist,
 dann R² H ist, R¹ F ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,
 f) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Ethylendioxyring bilden,
 dann R¹ Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,
 g) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Dihydrofuryl- oder Dihydropyrylring bilden,
 dann R¹ Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist, und
 h) wenn R² Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH oder SOMe ist,
 dann R¹ Et ist, R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

mit einem Ecdyson-Rezeptorkomplex innerhalb der Zellen der Pflanze, wobei die Zellen weiterhin eine DNA-Bindungssequenz für den Ecdyson-Rezeptorkomplex enthalten, wenn sie in Kombination mit dem Liganden vorliegen,

und wobei die Bildung eines Ecdyson-Rezeptorkomplex-Ligand-DNA-Bindungssequenz-Komplexes die Expression des Gens induziert.

5. Verfahren nach einem der Ansprüche 1 bis 4, wobei der Ligand die spezifizierte Formel hat und E t-Butyl ist.
6. Verfahren nach einem der Ansprüche 1 bis 5, wobei der Ligand die spezifizierte Formel hat und R¹ Me, Et, i-Pr oder F ist.
7. Verfahren nach einem der Ansprüche 1 bis 6, wobei der Ligand die spezifizierte Formel hat und R² OH, OMe, OEt ist oder verbunden ist mit R³ und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy- oder Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist.
8. Verfahren nach einem der Ansprüche 1 bis 7, wobei der Ligand die spezifizierte Formel hat und R³ H, Et ist oder verbunden ist mit R² und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy- oder Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist.
9. Verfahren nach einem der Ansprüche 1 bis 4, wobei der Ligand die spezifizierte Formel hat und E t-Butyl ist, R¹ Et ist, R² OEt ist, R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist.
10. Verfahren nach einem der Ansprüche 1 bis 9, wobei der Ecdyson-Rezeptorkomplex ein chimärer Ecdyson-Rezeptorkomplex ist und das DNA-Konstrukt weiterhin einen Promotor umfaßt.
11. Verwendung eines Liganden der Formel (1):



wobei:

- E ein (C₄-C₆)-Alkyl ist, welches einen tertiären Kohlenstoff enthält,
 R¹ H, Me, Et, i-Pr, F oder OMe ist,
 R² H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, S^{OMe} ist oder verbunden ist mit R³ und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,
 R³ H, Et ist oder verbunden ist mit R² und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,
 R⁴, R⁵ und R⁶ unabhängig voneinander H, Me, Et, F, Cl, CH₂OH, CN, OMe oder OEt sind, mit der Maßgabe, daß:
 a) wenn R¹ Me ist und R² OMe ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl oder 3,5-Di-F ist,
 b) wenn R¹ Me ist und R² OEt ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl ist,
 c) wenn R¹ Et ist und R² OMe oder OEt ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ folgendes ist:

- i) 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl oder 3-Me oder
- ii) R⁶ H ist, R⁴ Me ist und R⁵ Et, F, Cl, CH₂OH, CN, OMe oder OEt ist,

- d) wenn R¹ i-Pr ist,
dann R² OMe oder OEt ist, R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,
- e) wenn R³ Et ist,
dann R² H ist, R¹ F ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,
- f) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Ethylendioxyring bilden,
dann R¹ Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,
- g) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Dihydrofuryl- oder Dihydropyrylring bilden,
dann R¹ Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist, und
- h) wenn R² Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH oder SOMe ist,
dann R¹ Et ist, R³ H ist, die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

zum Modulieren der Expression eines oder mehrerer exogener Gene in einer Tierzelle *in vitro*, wobei die Zelle folgendes enthält:

- a) einen Ecdyson-Rezeptorkomplex, welcher folgendes umfaßt:

- 1) eine DNA-Bindungsdomäne,
- 2) eine Bindungsdomäne für den Liganden und
- 3) eine Transaktivierungsdomäne und

- b) ein DNA-Konstrukt, welches folgendes umfaßt:

- 1) das exogene Gen und
- 2) ein Antwortelement, und

wobei:

- a) das exogene Gen der Kontrolle durch das Antwortelement untersteht und
- b) eine Bindung der DNA-Bindungsdomäne an das Antwortelement in der Gegenwart des Liganden zur Aktivierung oder Unterdrückung des Gens führt.

12. Verwendung nach Anspruch 11, wobei der Ligand die spezifizierte Formel hat und E t-Butyl ist.

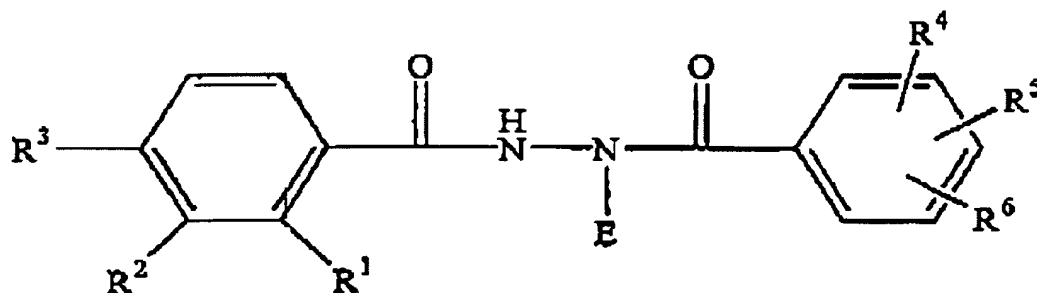
13. Verwendung nach Anspruch 11 oder 12, wobei der Ligand die spezifizierte Formel hat und R¹ Me, Et, i-Pr oder F ist.

14. Verwendung nach einem der Ansprüche 11 bis 13, wobei der Ligand die spezifizierte Formel hat und R² OH, OMe, OEt ist oder verbunden ist mit R³ und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy- oder Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist.

15. Verwendung nach einem der Ansprüche 11 bis 14, wobei der Ligand die spezifizierte Formel hat und R³ H, Et ist oder verbunden ist mit R² und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy- oder Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist.

16. Verwendung nach einem der Ansprüche 11 bis 15, wobei der Ligand die spezifizierte Formel hat und E t-Butyl ist, R¹ Et ist, R² OEt ist, R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist.

17. Verwendung eines Liganden der Formel (1):



wobei:

E ein (C₄-C₆)-Alkyl ist, welches einen tertiären Kohlenstoff enthält,
 R¹ H, Me, Et, i-Pr, F oder OMe ist,
 R² H, Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe ist oder verbunden ist mit R³ und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,
 R³ H, Et ist oder verbunden ist mit R² und den Phenylkohlenstoffen, an die R² und R³ angebunden sind, unter Bildung eines Ethylendioxy-, eines Dihydrofurylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist, oder eines Dihydropyrylrings, wobei der Sauerstoff zu einem Phenylkohlenstoff benachbart ist,
 R⁴, R⁵ und R⁶ unabhängig voneinander H, Me, Et, F, Cl, CH₂OH, CN, OMe oder OEt sind,

mit der Maßgabe, daß:

a) wenn R¹ Me ist und R² OMe ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl oder 3,5-Di-F ist,
 b) wenn R¹ Me ist und R² OEt ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me, 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl ist,
 c) wenn R¹ Et ist und R² OMe oder OEt ist,
 dann R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ folgendes ist:

i) 3,5-Di-OMe-4-Me, 3,5-Di-Cl, 3,5-Di-F, 2,4- oder 2,5-Di-F, 2,4- oder 2,5-Di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl oder 3-Me oder
 ii) R⁶ H ist, R⁴ Me ist und R⁵ Et, F, Cl, CH₂OH, CN, OMe oder OEt ist,

d) wenn R¹ i-Pr ist,
 dann R² OMe oder OEt ist, R³ H ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,
 e) wenn R³ Et ist,
 dann R² H ist, R¹ F ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,
 f) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Ethylendioxyring bilden,
 dann R¹ Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,
 g) wenn R² und R³ zusammen mit den Phenylkohlenstoffen, an die sie angebunden sind, einen Dihydrofuryl- oder Dihydropyrylring bilden,
 dann R¹ Et ist und die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist und
 h) wenn R² Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH oder SOMe ist,
 dann R¹ Et ist, R³ H ist, die Kombination aus R⁴, R⁵ und R⁶ 3,5-Di-Me ist,

bei der Herstellung eines Arzneimittels zum Modulieren der Expression eines oder mehrerer exogener Gene in einem Tier, wobei die Zellen des Tieres folgendes enthalten:

a) einen Ecdyson-Rezeptorkomplex, welcher folgendes umfaßt:

- 1) une DNA-Bindingsdomäne,
- 2) une Bindingsdomäne pour le Liganden und
- 3) une Transaktivierungsdomäne und

b) ein DNA-Konstrukt, welches folgendes umfaßt:

- 1) das exogene Gen und
- 2) ein Antwortelement, und

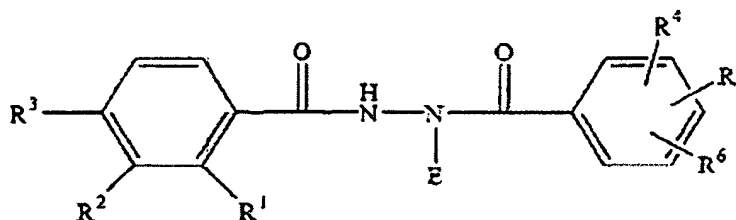
wobei:

- a) das exogene Gen der Kontrolle durch das Antwortelement untersteht und
- b) eine Bindung der DNA-Bindingsdomäne an das Antwortelement in der Gegenwart des Liganden zur Aktivierung oder Unterdrückung des Gens führt.

Revendications

1. Méthode *in vitro* pour moduler l'expression de gènes exogènes, comprenant la mise en contact d'un complexe de récepteur d'ecdysone comprenant :

- a) un domaine de liaison d'ADN ;
- b) un domaine de liaison de ligand ;
- c) un domaine de transactivation ; et
- d) un ligand de formule :



dans laquelle :

E représente un groupe alkyle en C₄ à C₆ contenant un atome de carbone tertiaire ;
 R¹ représente H, un groupe Me, Et, i-Pr, F ou OMe ;
 R² représente H, un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SMe, ou bien est joint avec R³ et les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;
 R³ représente H, un groupe Et, ou bien est joint avec R² et les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;
 R⁴, R⁵ et R⁶ représentent indépendamment H, un groupe Me, Et, F, Cl, CH₂OH, CN, OMe ou OEt ;
 sous réserve que :

- a) lorsque R¹ représente un groupe Me et R² représente un groupe OMe ;
 alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl ou 3, 5-di-F ;
- b) lorsque R¹ représente un groupe Me et R² représente un groupe OEt ;
 alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2, 5-di-Cl ;
- c) lorsque R¹ représente un groupe Et et R² représente un groupe OMe ou OEt ;

alors R^3 représente H ; et l'association de R^4 , R^5 et R^6 représente :

- i) un groupe 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl ou 3-Me ; ou
 ii) R^6 représente H, R^4 représente un groupe Me et R^5 représente un groupe Et, F, Cl, CH_2OH , CN, OMe ou OEt ;

d) lorsque R^1 représente un groupe i-Pr ;

alors R^2 représente un groupe OMe ou OEt ; R^3 représente H et l'association de R^4 , R^5 et R^6 représente un groupe 3,5-di-Me ;

e) lorsque R^3 représente un groupe Et ;

alors R^2 représente H ; R^1 représente F et l'association de R^4 , R^5 et R^6 représente un groupe 3,5-di-Me ;

f) lorsque R^2 et R^3 , conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau éthylènedioxy ;

alors R^1 représente un groupe Me ou Et et l'association de R^4 , R^5 et R^6 représente un groupe 3,5-di-Me ;

g) lorsque R^2 et R^3 , conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau dihydrofuryle ou dihydropyryle ;

alors R^1 représente un groupe Et et l'association de R^4 , R^5 et R^6 représente un groupe 3,5-di-Me ; et

h) lorsque R^2 représente un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH ou SOMe ;

alors R^1 représente un groupe Et, R^3 représente H et l'association de R^4 , R^5 et R^6 représente un groupe 3, 5-di-Me ;

avec un produit d'assemblage d'ADN comprenant :

a) le gène exogène ; et

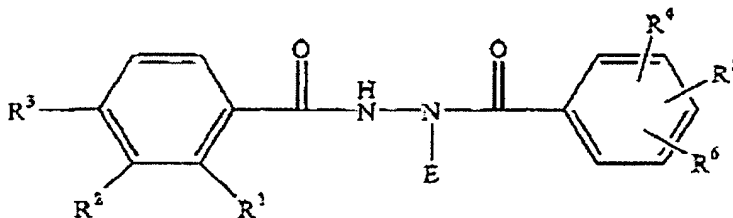
b) un élément de réponse ;

dans lequel :

a) le gène exogène est sous le contrôle de l'élément de réponse ; et

b) la liaison du domaine de liaison d'ADN à l'élément de réponse en présence du ligand a pour résultat l'activation ou la suppression du gène.

2. Procédé pour moduler l'expression d'un ou plusieurs gènes exogènes dans une plante, comprenant l'administration à la plante d'une quantité efficace d'un ligand de formule :



dans laquelle :

E représente un groupe alkyle en C_4 à C_6 contenant un atome de carbone tertiaire ;

R^1 représente H, un groupe Me, Et, i-Pr, F ou OMe ;

R^2 représente H, un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe, ou bien est joint avec R^3 et les atomes de carbone du groupe phényle auxquels R^2 et R^3 sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;

R^3 représente H, un groupe Et, ou bien est joint avec R^2 et les atomes de carbone du groupe phényle auxquels R^2 et R^3 sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;

R^4 , R^5 et R^6 représentent indépendamment H, un groupe Me, Et, F, Cl, CH_2OH , CN, OMe ou OEt ;

sous réserve que :

a) lorsque R¹ représente un groupe Me et R² représente un groupe OMe ;
alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl ou 3,5-di-F ;

b) lorsque R¹ représente un groupe Me et R² représente un groupe OEt ;
alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl ;

c) lorsque R¹ représente un groupe Et et R² représente un groupe OMe ou OEt ;
alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente :

i) un groupe 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl ou 3-Me ; ou

ii) R⁶ représente H, R⁴ représente un groupe Me et R⁵ représente un groupe Et, F, Cl, CH₂OH CN, OMe ou OEt ;

d) lorsque R¹ représente un groupe i-Pr ;
alors R² représente un groupe OMe ou OEt ; R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

e) lorsque R³ représente un groupe Et ;

alors R² représente H ; R¹ représente F et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

f) lorsque R² et R³, conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau éthylènedioxy ;

alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

g) lorsque R² et R³, conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau dihydrofuryle ou dihydropyryle ;

alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ; et

h) lorsque R² représente un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH ou SOMe ;

alors R¹ représente un groupe Et, R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

dans lequel les cellules de la plante contiennent :

a) un complexe de récepteur d'ecdysone comprenant :

1) un domaine de liaison d'ADN ;

2) un domaine de liaison pour le ligand ; et

3) un domaine de transactivation ; et

b) un produit d'assemblage d'ADN comprenant :

1) le gène exogène ; et

2) un élément de réponse ; et

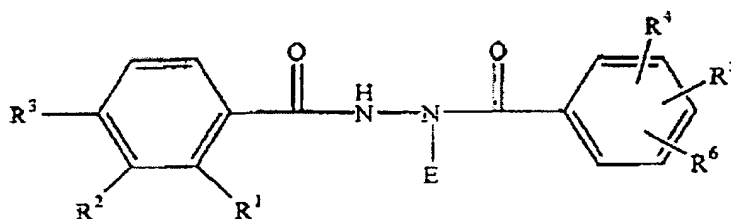
dans lequel :

a) le gène exogène est sous le contrôle de l'élément de réponse ; et

b) la liaison du domaine de liaison d'ADN à l'élément de réponse en présence du ligand a pour résultat l'activation ou la suppression du gène.

3. Procédé pour la production d'un polypeptide, comprenant les étapes consistant à :

a) sélectionner une cellule qui est insensible à l'exposition à un ligand de formule :



dans laquelle :

E représente un groupe alkyle en C₄ à C₆ contenant un atome de carbone tertiaire ;

R¹ représente H, un groupe Me, Et, i-Pr, F ou OMe ;

R² représente H, un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe, ou bien est joint avec R³ et les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;

R³ représente H, un groupe Et, ou bien est joint avec R² et les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;

R⁴, R⁵ et R⁶ représentent indépendamment H, un groupe Me, Et, F, Cl, CH₂OH, CN, OMe ou OEt ;

sous réserve que :

a) lorsque R¹ représente un groupe Me et R² représente un groupe OMe ;

alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl ou 3,5-di-F ;

b) lorsque R¹ représente un groupe Me et R² représente un groupe OEt ;

alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl ;

c) lorsque R¹ représente un groupe Et et R² représente un groupe OMe ou OEt ;

alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente :

i) un groupe 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl ou 3-Me ; ou

ii) R⁶ représente H, R⁴ représente un groupe Me et R⁵ représente un groupe Et, F, Cl, CH₂OH, OMe ou OEt ;

d) lorsque R¹ représente un groupe i-Pr ;

alors R² représente un groupe OMe ou OEt ; R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

e) lorsque R³ représente un groupe Et ;

alors R² représente H ; R¹ représente F et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

f) lorsque R² et R³, conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau éthylènedioxy ;

alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

g) lorsque R² et R³, conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau dihydrofuryle ou dihydropyryle ;

alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ; et

h) lorsque R² représente un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH ou SOMe ;

alors R¹ représente un groupe Et, R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

b) à introduire dans la cellule :

1) un produit d'assemblage d'ADN comprenant :

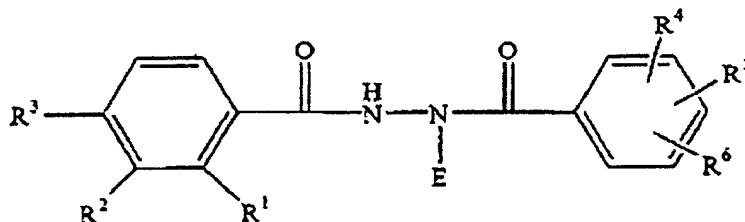
- a) un gène exogène codant pour le polypeptide, et
- b) un élément de réponse ;
- le gène étant sous le contrôle de l'élément de réponse ; et

2) un complexe de récepteur d'ecdysone comprenant .

- a) un domaine de liaison d'ADN ;
- b) un domaine de liaison pour le ligand ; et
- c) un domaine de transactivation ; et

c) à exposer la cellule au ligand de l'étape (a).

4. Procédé pour réguler l'expression de gènes endogènes ou hétérologues dans une plante transgénique, comprenant la mise en contact d'un ligand de formule :



dans laquelle :

E représente un groupe alkyle en C₄ à C₆ contenant un atome de carbone tertiaire ;

R¹ représente H, un groupe Me, Et, i-Pr, F ou OMe ;

R² représente H, un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SMe, ou bien est joint avec R³ et les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;

R³ représente H, un groupe Et, ou bien est joint avec R² et les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;

R⁴, R⁵ et R⁶ représentent indépendamment H, un groupe Me, Et, F, Cl, CH₂OH, CN, OMe ou OEt ;

sous réserve que :

a) lorsque R¹ représente un groupe Me et R² représente un groupe OMe ;

alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl ou 3,5-di-F ;

b) lorsque R¹ représente un groupe Me et R² représente un groupe OEt ;

alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl ;

c) lorsque R¹ représente un groupe Et et R² représente un groupe OMe ou OEt ;

alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente :

i) un groupe 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl ou 3-Me ; ou

ii) R⁶ représente H, R⁴ représente un groupe Me et R⁵ représente un groupe Et, F, Cl, CH₂OH, CN, OMe ou OEt ;

d) lorsque R¹ représente un groupe i-Pr ;

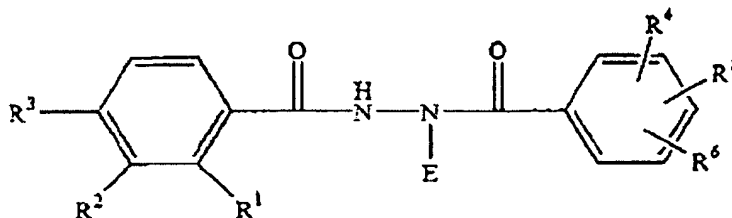
alors R² représente un groupe OMe ou OEt ; R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

e) lorsque R³ représente un groupe Et ;

alors R² représente H ; R¹ représente F et l'association de R⁴, R⁵ et R⁶ représente un groupe 3, 5-di-Me ;
 f) lorsque R² et R³ conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés,
 forment un noyau éthylènedioxy ;
 alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3, 5-di-Me ;
 g) lorsque R² et R³, conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés,
 forment un noyau dihydrofuryle ou dihydropyryle ;
 alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3, 5-di-Me ; et
 h) lorsque R² représente un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH ou SOMe ;
 alors R¹ représente un groupe Et, R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe
 3,5-di-Me ;

avec un complexe de récepteur d'ecdysone à l'intérieur des cellules de la plante, dans lequel les cellules contiennent
 en outre une séquence de liaison d'ADN pour le complexe de récepteur d'ecdysone lors de l'association avec le
 ligand et dans lequel la formation d'un complexe de récepteur d'ecdysone-ligand-séquence de liaison
 d'ADN induit l'expression du gène.

5. Procédé suivant l'une quelconque des revendications 1 à 4, dans lequel le ligand répond à la formule spécifiée et
 E représente un groupe tertibutyle.
6. Procédé suivant l'une quelconque des revendications 1 à 5, dans lequel le ligand répond à la formule spécifiée et
 R¹ représente un groupe Me, Et, i-Pr ou F.
7. Procédé suivant l'une quelconque des revendications 1 à 6, dans lequel le ligand répond à la formule spécifiée et
 R² représente un groupe OH, OMe, OEt, ou bien est joint avec R³ et avec les atomes de carbone du groupe phényle
 auxquels R² et R³ sont fixés pour former un noyau éthylènedioxy ou dihydrofuryle avec l'atome d'oxygène adjacent
 à un atome de carbone du groupe phényle.
8. Procédé suivant l'une quelconque des revendications 1 à 7, dans lequel le ligand répond à la formule spécifiée et
 R³ représente H ou un groupe Et, ou bien est joint avec R² et avec les atomes de carbone du groupe phényle
 auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy ou un noyau dihydrofuryle avec l'atome d'oxygène
 adjacent à un atome de carbone du groupe phényle.
9. Procédé suivant l'une quelconque des revendications 1 à 4, dans lequel le ligand répond à la formule spécifiée et
 E représente un groupe tertibutyle, R¹ représente un groupe Et, R² représente un groupe OEt, R³ représente H,
 et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me.
10. Procédé suivant l'une quelconque des revendications 1 à 9, dans lequel le complexe de récepteur d'ecdysone est
 un complexe de récepteur d'ecdysone chimère et le produit d'assemblage d'ADN comprend en outre un promoteur.
11. Utilisation d'un ligand de formule (I) :



dans laquelle :

E représente un groupe alkyle en C₄ à C₆ contenant un atome de carbone tertiaire ;
 R¹ représente H, un groupe Me, Et, i-Pr, F ou OMe ;
 R² représente H, un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe, ou bien est joint avec R³ et les
 atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un

noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;

R³ représente H, un groupe Et, ou bien est joint avec R² et les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle;

R⁴, R⁵ et R⁶ représentent indépendamment H, un groupe Me, Et, F, Cl, CH₂OH, CN, OMe ou OEt; sous réserve que :

a) lorsque R¹ représente un groupe Me et R² représente un groupe OMe ;
alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl ou 3,5-di-F ;

b) lorsque R¹ représente un groupe Me et R² représente un groupe OEt ;
alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl ;

c) lorsque R¹ représente un groupe Et et R² représente un groupe OMe ou OEt ;
alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente :

i) un groupe 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl ou 3-Me ; ou

ii) R⁶ représente H, R⁴ représente un groupe Me et R⁵ représente un groupe Et, F, Cl, CH₂OH, CN, OMe ou OEt ;

d) lorsque R¹ représente un groupe i-Pr ;
alors R² représente un groupe OMe ou OEt; R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me;

e) lorsque R³ représente un groupe Et;

alors R² représente H; R¹ représente F et l'association de R⁴, R⁵ et R⁶ représente un groupe 3, 5-di-Me;

f) lorsque R² et R³, conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau éthylènedioxy ;

alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

g) lorsque R² et R³, conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau dihydrofuryle ou dihydropyryle;

alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3, 5-di-Me ; et

h) lorsque R² représente un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH ou SOME;

alors R¹ représente un groupe Et, R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me;

pour la modulation *in vitro* de l'expression d'un ou plusieurs gènes exogènes dans une cellule animale, dans laquelle la cellule contient :

a) un complexe de récepteur d'ecdysone comprenant :

1) un domaine de liaison d'ADN ;

2) un domaine de liaison pour le ligand ; et

3) un domaine de transactivation; et

b) un produit d'assemblage d'ADN comprenant :

1) le gène exogène ; et

2) un élément de réponse ; et

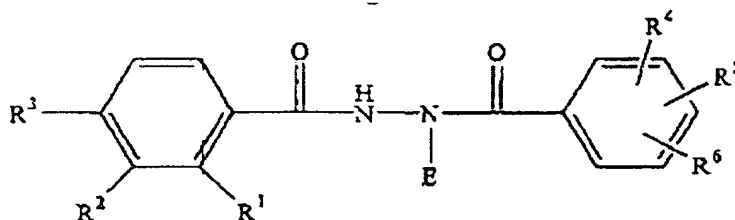
dans laquelle :

a) le gène exogène est sous le contrôle de l'élément de réponse ; et

b) la liaison du domaine de liaison d'ADN à l'élément de réponse en présence du ligand a pour résultat l'activation ou la suppression du gène.

12. Utilisation suivant la revendication 11, dans laquelle le ligand répond à la formule spécifiée et E représente un groupe tertibutyle.

13. Utilisation suivant la revendication 11 ou 12, dans laquelle le ligand répond à la formule spécifiée et R¹ représente un groupe Me, Et, i-Pr ou F.
14. Utilisation suivant l'une quelconque des revendications 11 à 13, dans laquelle le ligand répond à la formule spécifiée et R² représente un groupe OH, OMe, OEt, ou bien est joint avec R³ et avec les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un noyau éthylènedioxy ou dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle.
15. Utilisation suivant l'une quelconque des revendications 11 à 14, dans laquelle le ligand répond à la formule spécifiée et R³ représente H ou un groupe Et, ou bien est joint avec R² et avec les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un noyau éthylènedioxy ou dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle.
16. Utilisation suivant l'une quelconque des revendications 11 à 15, dans laquelle le ligand répond à la formule spécifiée et E représente un groupe tertibutyle, R¹ représente un groupe Et, R² représente un groupe OEt, R³ représente H, et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me.
17. Utilisation d'un ligand de formule (I) :



dans laquelle :

E représente un groupe alkyle en C₄ à C₆ contenant un atome de carbone tertiaire ;
 R¹ représente H, un groupe Me, Et, i-Pr, F ou OMe ;
 R² représente H, un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH, OMe, OEt, SOMe, ou bien est joint avec R³ et les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;
 R³ représente H, un groupe Et, ou bien est joint avec R² et les atomes de carbone du groupe phényle auxquels R² et R³ sont fixés pour former un groupe éthylènedioxy, un noyau dihydrofuryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle, ou un noyau dihydropyryle avec l'atome d'oxygène adjacent à un atome de carbone du groupe phényle ;
 R⁴, R⁵ et R⁶ représentent indépendamment H, un groupe Me, Et, F, Cl, CH₂OH, CN, OMe ou OEt ;
 sous réserve que :

- a) lorsque R¹ représente un groupe Me et R² représente un groupe OMe ;
 alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl ou 3, 5-di-F ;
 b) lorsque R¹ représente un groupe Me et R² représente un groupe OEt ;
 alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me, 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl ;
 c) lorsque R¹ représente un groupe Et et R² représente un groupe OMe ou OEt ;
 alors R³ représente H ; et l'association de R⁴, R⁵ et R⁶ représente :
 i) un groupe 3,5-di-OMe-4-Me, 3,5-di-Cl, 3,5-di-F, 2,4- ou 2,5-di-F, 2,4- ou 2,5-di-Cl, 3-OMe, 2-Cl-5-Me, 2-Cl ou 3-Me ; ou
 ii) R⁶ représente H, R⁴ représente un groupe Me et R⁵ représente un groupe Et, F, Cl, CH₂OH CN, OMe ou OEt ;
 d) lorsque R¹ représente un groupe i-Pr ;

alors R² représente un groupe OMe ou OEt ; R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

e) lorsque R³ représente un groupe Et ;

alors R² représente H ; R¹ représente F et l'association de R⁴, R⁵ et R⁶ représente un groupe 3, 5-di-Me ;

f) lorsque R² et R³, conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau éthylènedioxy ;

alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

g) lorsque R² et R³, conjointement avec les atomes de carbone du groupe phényle auxquels ils sont fixés, forment un noyau dihydrofuryle ou dihydropyryle ;

alors R¹ représente un groupe Et et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ; et

h) lorsque R² représente un groupe Me, Et, n-Pr, i-Pr, Ac, F, Cl, OH ou SMe ;

alors R¹ représente un groupe Et, R³ représente H et l'association de R⁴, R⁵ et R⁶ représente un groupe 3,5-di-Me ;

dans la préparation d'un agent pharmaceutique pour la modulation de l'expression d'un ou plusieurs gènes exogènes chez un animal, dans laquelle les cellules de l'animal contiennent :

a) un complexe de récepteur d'ecdysone comprenant :

1) un domaine de liaison d'ADN ;

2) un domaine de liaison pour le ligand ; et

3) un domaine de transactivation ; et

b) un produit d'assemblage d'ADN comprenant :

1) le gène exogène ; et

2) un élément de réponse ; et

dans laquelle :

a) le gène exogène est sous le contrôle de l'élément de réponse ; et

b) la liaison du domaine de liaison d'ADN à l'élément de réponse en présence du ligand a pour résultat l'activation ou la suppression du gène.

REFERENCES CITED IN THE DESCRIPTION

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